

19



Europäisches Patentamt
European Patent Office
Office européen des brevets

11 Publication number:

**0 379 055
A2**

12

EUROPEAN PATENT APPLICATION

21 Application number: 90100485.3

51 Int. Cl.⁵: C08G 59/02, C08G 59/20

22 Date of filing: 11.01.90

30 Priority: 17.01.89 US 297896

43 Date of publication of application:
25.07.90 Bulletin 90/30

84 Designated Contracting States:
AT BE CH DE ES FR GB GR IT LI NL SE

71 Applicant: THE DOW CHEMICAL COMPANY
2030 Dow Center Abbott Road P.O. Box 1967
Midland Michigan 48640-1967(US)

72 Inventor: Earls, Jimmy D.
209 Banyan
Lake Jackson Texas 77566(US)
Inventor: Puckett, Paul M.
126 Daisy
Lake Jackson Texas 77566(US)

74 Representative: Sternagel, Hans-Günther, Dr.
et al
Patentanwälte Dr. Michael Hann Dr. H.-G.
Sternagel Sander Aue 30
D-5060 Bergisch Gladbach 2(DE)

54 Mesogenic advanced epoxy compounds.

57 Advanced epoxy resin compounds are prepared by reacting weight epoxy resins with active hydrogen-containing compounds which contain rigid, mesogenic type structures.

EP 0 379 055 A2

MESOGENIC ADVANCED EPOXY COMPOUNDS

The present invention concerns advanced epoxy resin compounds containing mesogenic moieties, curable compositions and cured compositions thereof.

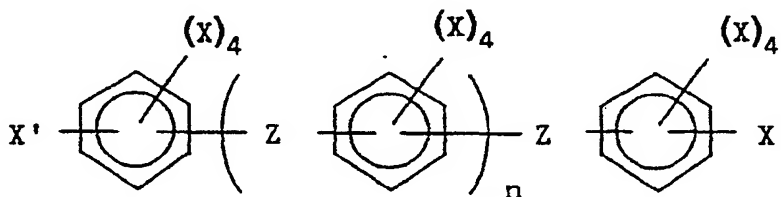
Epoxy resins are useful in many applications such as adhesives, coatings, castings, laminates, and composites. In each of these applications, it is desirable to have epoxy resins with improved physical and/or thermal properties.

The present invention provides advanced epoxy resin compounds with improved properties through the incorporation of mesogenic (liquid crystalline type) structures into the polymer chain. These resins are obtained through advancement reactions with mesogenic compounds containing active hydrogens. The typical structure of the mesogenic compounds consists of two or more aromatic rings bridged by a rigid, central linkage. The active hydrogen sites are provided by hydroxyl, amine, amide and/or carboxylic acid functionality.

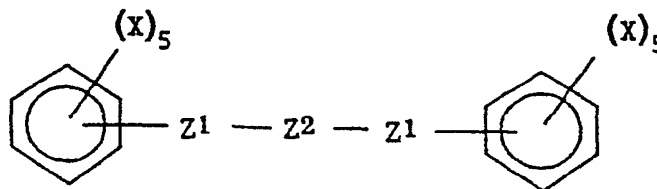
An additional aspect of the present invention is that the property improvements which are obtained with these resins can be further enhanced by the application of electric and magnetic fields and shear stresses during processing and/or cure.

One aspect of the present invention pertains to advanced epoxy resin compounds prepared by reacting an epoxy resin having an average of more than one vicinal epoxy group per molecule with a compound having an average of more than one active hydrogen atom per molecule represented by the following formulas I, II or III:

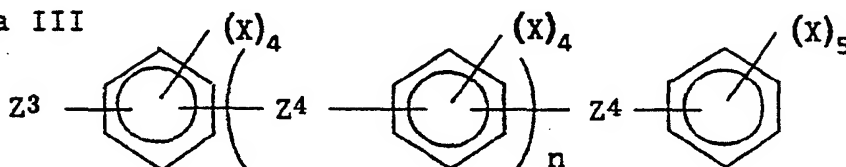
Formula I



Formula II

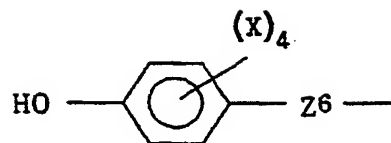


Formula III

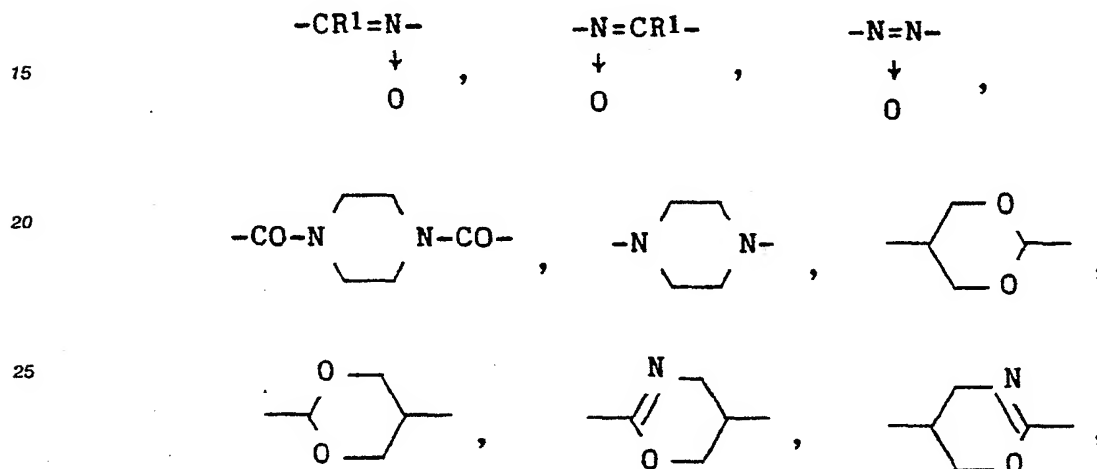


wherein at least about 80 percent of the Z and X' groups are in the para position with respect to each other in Formula I; at least about 80 percent of the Z³ and Z⁴ groups are in the para position with respect to each other in Formula III; each X is independently hydrogen, a hydrocarbyl or hydrocarbyloxy group having from 1 to 12, preferably from 1 to 6, most preferably from 1 to 4, carbon atoms, a halogen atom, -NO₂ or -C≡N; each X' is independently a hydroxyl group, a carboxylic acid group or the group represented by formula IV:

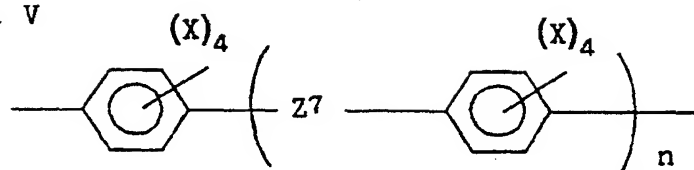
Formula IV



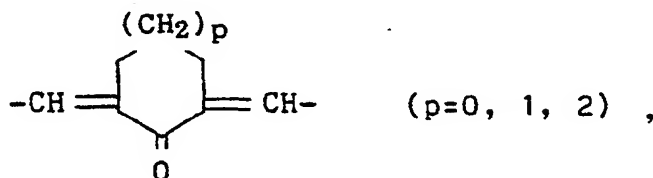
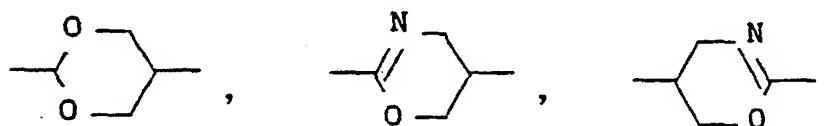
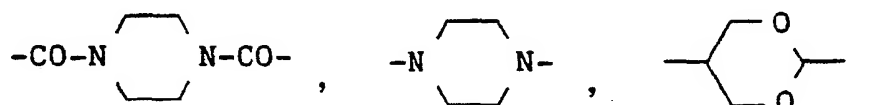
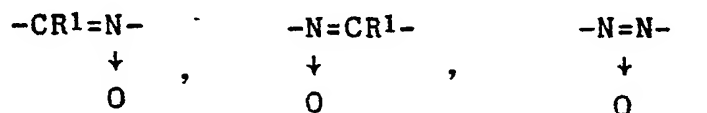
each Z is independently $-\text{CR}^1 = \text{CR}^1-$, $-\text{CR}^1 = \text{CR}^1 - \text{CR}^1 = \text{CR}^1-$, $-\text{CR}^1 = \text{N} - \text{N} = \text{CR}^1-$, $-\text{CR}^1 = \text{CR}^1 - \text{CO} - \text{O} - \text{CH}_2-$,
 10 $-\text{CR}^1 = \text{CR}^1 - \text{CO} - \text{O} - \text{CH}_2 - \text{CH}_2-$, $-\text{CH}_2 - \text{O} - \text{CO} - \text{CR}^1 = \text{CR}^1-$, $-\text{CH}_2 - \text{CH}_2 - \text{O} - \text{CO} - \text{CR}^1 = \text{CR}^1-$, $-\text{CR}^1 = \text{CR}^1 - \text{CO} - \text{O}-$, $-\text{O} -$
 $\text{CO} - \text{CR}^1 = \text{CR}^1-$, $-\text{N} = \text{N}-$, $-\text{CO} - \text{NH}-$, $-\text{NH} - \text{CO}-$, $-\text{CO} - \text{NH} - \text{NH} - \text{CO}-$, $-\text{C} \equiv \text{C}-$, $-\text{C} \equiv \text{C} - \text{C} \equiv \text{C}-$, $-\text{CO} - \text{S}-$, $-\text{S} - \text{CO}-$, $-\text{CO} - \text{O}-$,
 $-\text{O} - \text{CO}-$, a direct single bond when $n \geq 1$,



Formula V



Z^3 is $\text{NH}_2\text{---}$, $\text{NH}_2\text{---SO}_2\text{---}$, $\text{NH}_2\text{---CO---}$, or $\text{NH}_2\text{---Z}^5\text{---O---}$; each Z^4 is independently $\text{---CR}^1=\text{CR}^1\text{---}$, $\text{---CR}^1=\text{CR}^1\text{---}$, $\text{---CR}^1=\text{N---N---CR}^1\text{---}$, $\text{---CR}^1=\text{CR}^1\text{---CO---O---CH}_2\text{---}$, $\text{---CR}^1=\text{CR}^1\text{---CO---O---CH}_2\text{---CH}_2\text{---}$, $\text{---CH}_2\text{---O---CO---CR}^1=\text{CR}^1\text{---}$, $\text{---CH}_2\text{---CH}_2\text{---O---CO---CR}^1=\text{CR}^1\text{---}$, $\text{---CR}^1=\text{CR}^1\text{---CO---O---}$, $\text{---O---CO---CR}^1=\text{CR}^1\text{---}$, ---N=N--- , ---CO---NH--- , ---NH---CO--- , ---CO---NH--- , ---NH---CO--- , $\text{---C}\equiv\text{C---}$, $\text{---C}\equiv\text{C---C}\equiv\text{C---}$, ---CO---S--- , ---S---CO--- , $\text{---CR}^1=\text{N---}$, $\text{---N=CR}^1\text{---}$, ---CO---O--- , ---O---CO--- , a direct single bond,



5

10

15

20

25

30

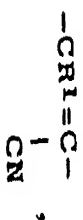
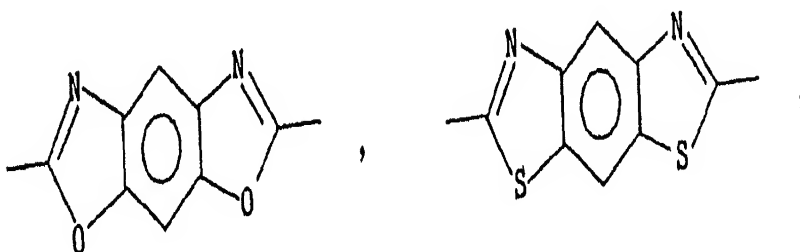
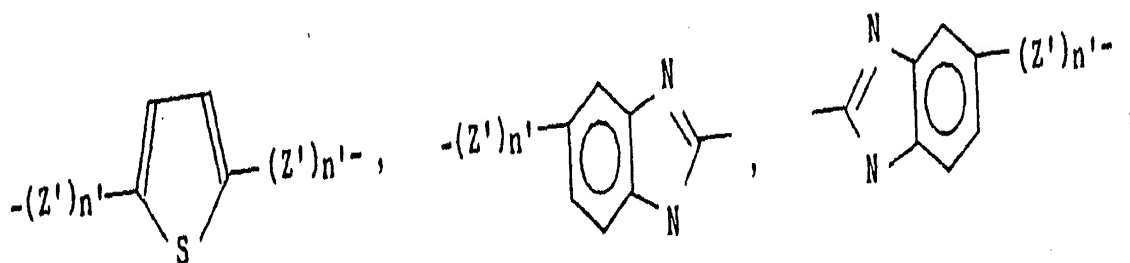
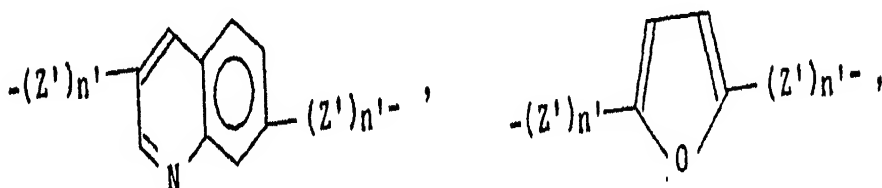
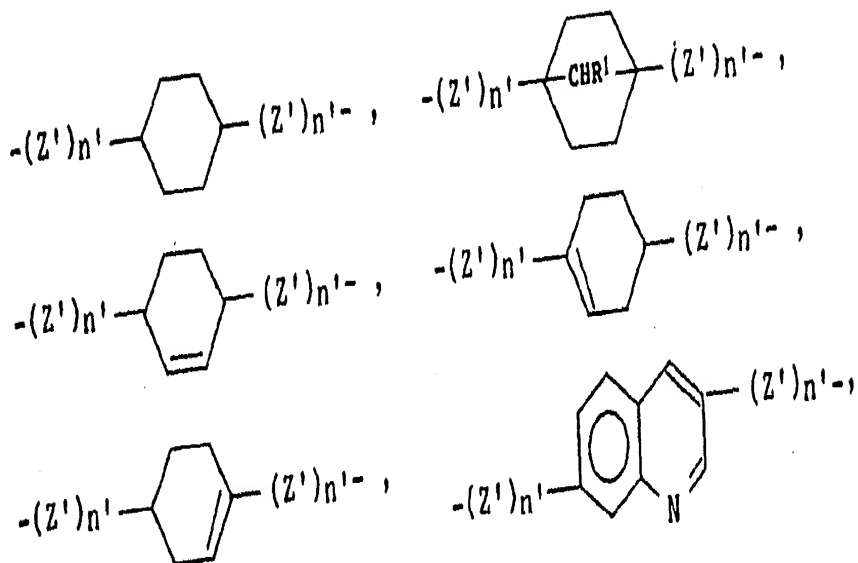
35

40

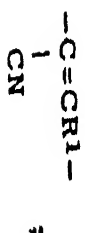
45

50

55



and



Z⁵ is an alkyl or cycloalkyl group having from 1 to 10, preferably from 1 to 6, more preferably from 1 to 3 carbon atoms; Z⁷ is the same as Z⁴ with the proviso that it can also independently be a divalent hydrocarbyl group having from 1 to 4, carbon atoms, -SO-, -SO₂-, -S-, -S-S-, -O-, or -CO-; and n has an average value of zero to 6.

5 Another aspect of the present invention pertains to curable compositions comprising the aforementioned advanced epoxy resins and a curing amount of a curing agent or curing catalyst therefor.

A further aspect of the present invention pertains to products or articles resulting from curing the aforementioned curable compositions.

10 A still further aspect of the present invention pertains to products resulting from the application of an electric field or magnetic field or shear flow during curing or processing of the aforementioned advanced epoxy compounds.

The advanced epoxy resin compounds of the present invention provide cured products having an improvement in one or more of the following physical or thermal properties such as, but not limited to, glass transition temperature, tensile strength, tensile modulus, flexural strength, and flexural modulus.

15 The term "hydrocarbyl" as employed herein means any aliphatic, cycloaliphatic, aromatic, aryl substituted aliphatic or cycloaliphatic, or aliphatic or cycloaliphatic substituted aromatic groups. The aliphatic groups can be saturated or unsaturated. Likewise, the term "hydrocarbyloxy" means a hydrocarbyl group having an oxygen linkage between it and the carbon atom to which it is attached.

20 The term "active hydrogen atom" as employed herein means a hydrogen atom which is reactive with a vicinal epoxide group.

Particularly suitable compounds having an average of more than one active hydrogen atom per molecule which can be employed herein include hydroxyl-containing compounds, primary amine-containing compounds and compounds containing more than one aromatic amide group per molecule. These compounds include, for example, those compounds represented by the aforementioned formulas I, II and III.

25 Particularly suitable hydroxyl-containing compounds include, for example, bis(4-hydroxyphenyl)-terephthalate, N,N'-bis(4-hydroxyphenyl)terephthalamide, bis(4-hydroxybiphenyl)terephthalate, 4,4'-dihydroxyphenylbenzoate, 4,4'-dihydroxybenzanilide, 4,4'-dihydroxybiphenylbenzoate, 1,4-bis(4'-hydroxyphenyl-1'-carboxamide)benzene, 1,4-bis(4'-hydroxyphenyl-1'-carboxy)benzene, 4,4'-bis(4''-hydroxyphenyl-1'-carboxy)biphenyl, bis(4'-hydroxyphenyl)-1,4-benzenediimine, 4,4'-dihydroxy- α -methylstilbene, 4,4'-dihydroxy- α -cyanostilbene, 4,4'-dihydroxystilbene, 4,4'-dihydroxy-2,2'-dimethylazoxybenzene, 4,4'-dihydroxy- α,α -diethylstilbene and mixtures thereof.

Particularly suitable carboxylic acid-containing compounds include, for example, 4,4'-benzanilide dicarboxylic acid, 4,4'-phenylbenzoate dicarboxylic acid, 4,4'-stilbenedicarboxylic acid and mixtures thereof.

35 Particularly suitable primary amine-containing compounds include, for example, 4'-sulfonamido-N-phenyl benzamide, 4'-sulfonamido-N'-phenyl-4-chlorobenzamide, 4-amino-1-phenylbenzoate, 4-amino-N-phenylbenzamide, N-phenyl-4-aminophenyl-1-carboxamide, phenyl-4-aminobenzoate, biphenyl-4-aminobenzoate, 1-phenyl-4'-aminophenylterephthalate, and mixtures thereof.

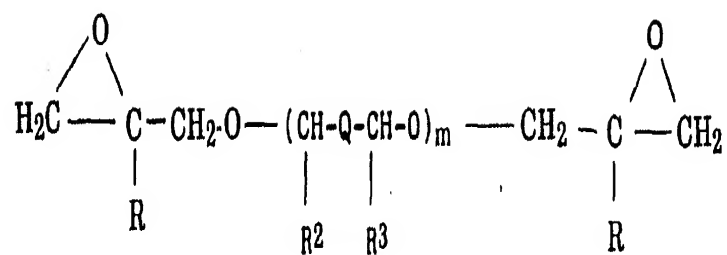
40 Particularly suitable aromatic amide-containing compounds include, for example, N,N'-diphenylterephthalamide, 1,4-benzenediphenylcarboxamide, 1,4-cyclohexanediphenylcarboxamide, benzanilide ether, N-phenylbenzamide and mixtures thereof.

Suitable epoxy resins which can be advanced with the aforementioned active hydrogen-containing compounds include those epoxy resins having an aliphatic, cycloaliphatic or aromatic base. These include the polyglycidyl ethers of alkylene glycols, polyoxyalkylene glycols, dihydric phenols including bisphenols and dihydroxy biphenyl as well as substituted derivatives thereof, phenol and substituted phenol aldehyde 45 novolac resins, phenol or substituted phenol hydrocarbon resins. The dihydric phenols and novolac resins and hydrocarbon resins can contain such substituents as hydrocarbyl or hydrocarbyloxy groups containing from 1 to 10, preferably from 1 to 6, more preferably from 1 to 3 carbon atoms, halogen atoms, particularly chlorine or bromine, -NO₂, and -C \equiv N. Exemplary of such suitable epoxy resins include those represented by the following formulas VI, VII, VIII, IX or X:

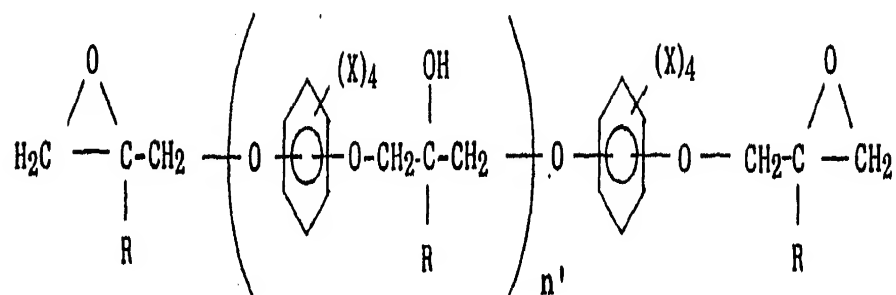
50

55

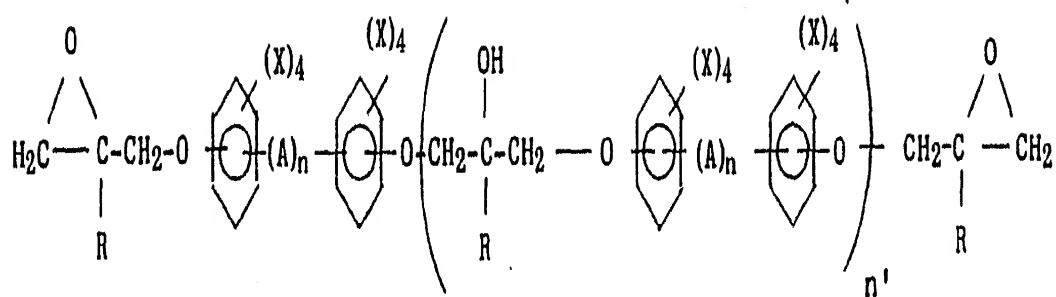
Formula VI



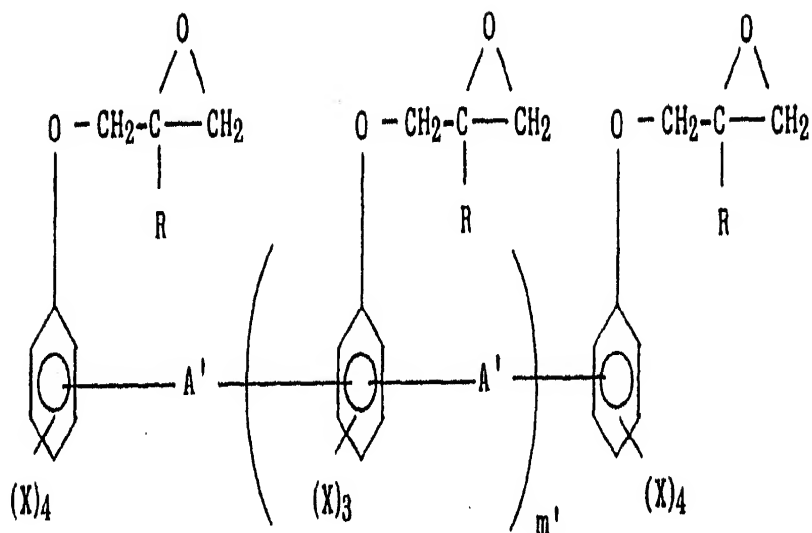
FORMULA VII



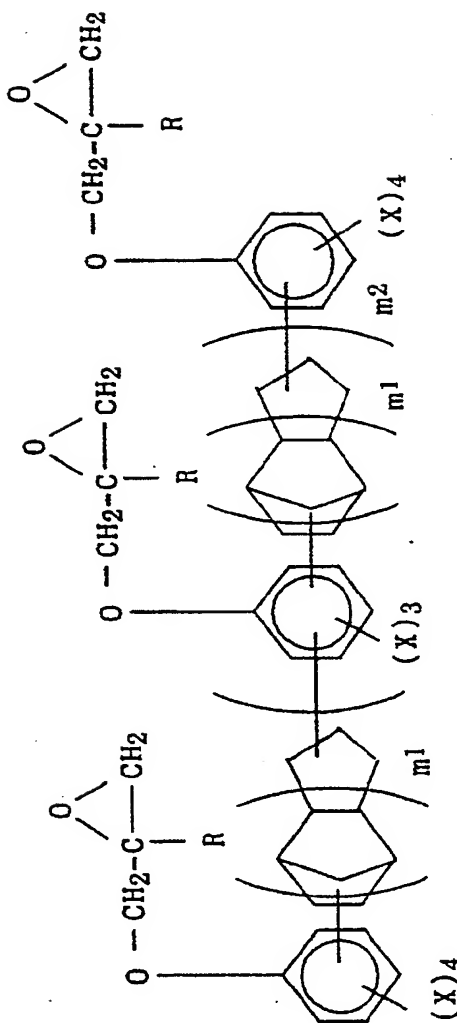
FORMULA VIII



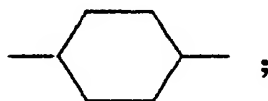
FORMULA IX



FORMULA X



wherein each A is independently a divalent hydrocarbyl group having from 1 to 12, preferably from 1 to 6, more preferably from 1 to 3 carbon atoms, -O-, -S-, -S-S-, -SO-, -SO₂-, or -CO-; each A' independently a divalent hydrocarbon group having from 1 to 6, preferably from 1 to 3 carbon atoms; Q is a single bond, -CH₂-S-CH₂-, -(CH₂)_n-, or



each R is independently hydrogen or an alkyl group having from 1 to 4 carbon atoms; each R² and R³ is independently hydrogen, a hydrocarbyl or halohydrocarbyl group having from 1 to 6, preferably from 1 to 3, more preferably from 1 to 2 carbon atoms; each X is independently hydrogen, a hydrocarbyl or hydrocarbyloxy group having from 1 to 12, preferably from 1 to 6, most preferably from 1 to 4 carbon atoms, a halogen atom, -NO₂ or -C≡N; m has a value from 1 to 10, preferably from 1 to 4, more preferably from 1 to 2; m' has a value from 0.01 to 12, preferably from 1 to 6, more preferably from 1 to 3; m¹ has an average value from 1 to 12, preferably from 1 to 6, more preferably from 1 to 3; m² has a value from 1 to 12, preferably from 2 to 6, more preferably from 2 to 3; n has a value of zero or 1; n' has an average value from zero to 3, preferably from zero to 1.5, more preferably from zero to 0.5; and n¹ has an average value from 1 to 10.

The epoxy resin having an average of more than one vicinal epoxy group per molecule and the

compound having an average of more than one active hydrogen atom per molecule are reacted in amounts which provide suitably from 0.01:1 to 0.05:1, more suitably from 0.01:1 to 0.5:1, most suitably from 0.01:1 to 0.3:1 active hydrogen atoms per epoxy group.

When the epoxy resin having an average of more than one vicinal epoxy group per molecule and the compound having an average of more than one active hydrogen atom per molecule and containing one or more mesogenic structures are reacted in amounts which provide from 0.01:1 to 0.95:1 active hydrogen atoms per epoxy group, a curable (thermosettable) advanced epoxy resin is produced. When the epoxy resin having an average of more than one vicinal epoxy group per molecule and the compound having an average of more than one active hydrogen atom per molecule and containing one or more mesogenic structures are reacted in amounts which provide from 0.96:1 to 1.05:1 active hydrogen atoms per epoxy group, a substantially thermoplastic, resinous product is provided. Said resin composition contains little, if any, curable residual epoxide functionality, and may thus be processed using the typical processing methods employed with conventional thermoplastics, such as, for example, injection molding or extrusion. Thermosetting may, however, be induced, for example, via reaction of all or a part of the backbone secondary hydroxyl groups produced in the aforesaid advancement reaction, with a curing agent therefor. One class of said curing agents includes the di/polyisocyanates as well as the blocked di/polyisocyanates which can be induced to react with said secondary hydroxy groups providing urethane linked crosslinks between the resin chains. An example of a specific diisocyanate especially useful herein is 4,4'-diisocyanatodiphenylmethane. When the compounds having an average of more than one active hydrogen atom per molecule used in the advancement reaction is a diphenol, the resultant resinous product is a phenoxy resin. According to the teachings found in Encyclopedia of Polymer Science and Engineering, Vol. 6, page 331, published by John Wiley and Sons, New York (1986) aside from the aforementioned advancement method, a phenoxy resin may also be prepared by reaction of a 1:1 mole ratio of high purity bisphenol A and epichlorohydrin. It is therefore operable to prepare the phenoxy resins containing one or more mesogenic structures of the present invention via reaction of one or more diphenols containing one or more mesogenic structures with one or more epichlorohydrins. A typical example would thus be the phenoxy resin produced from the reaction of epichlorohydrin and bis(4-hydroxyphenyl)terephthalate using the aforementioned stoichiometric ratio.

The advancement reaction can be conducted in the presence of a suitable advancement catalyst such as phosphines, quaternary ammonium compounds, phosphonium compounds and tertiary amines. Particularly suitable catalysts include, for example, ethyltriphenylphosphonium chloride, ethyltriphenylphosphonium bromide, ethyltriphenylphosphonium iodide, ethyltriphenylphosphonium diacetate (ethyltriphenylphosphonium acetate*acetic acid complex), ethyltriphenylphosphonium phosphate, tetrabutylphosphonium chloride, tetrabutylphosphonium bromide, tetrabutylphosphonium iodide, tetrabutylphosphonium diacetate (tetrabutylphosphonium acetate*acetic acid complex), butyltriphenylphosphonium tetrabromobisphenate, butyltriphenylphosphonium bisphenate, butyltriphenylphosphonium bicarbonate, benzyltrimethylammonium chloride, tetramethylammonium hydroxide, triethylamine, tripropylamine, tributylamine, 2-methylimidazole, benzyltrimethylamine and mixtures thereof. Many of these catalysts are described in U. S. Patent Nos. 3,306,872; 3,341,580; 3,379,684; 3,477,990; 3,547,881; 3,637,590; 3,843,605; 3,948,855; 3,956,237; 4,048,141; 4,093,650; 4,131,633; 4,132,706; 4,171,420; 4,177,216 and 4,366,295.

The amount of advancement catalyst depends, of course, upon the particular reactants and catalyst employed; however, the advancement catalyst is usually employed in quantities of from 0.03 to 3, preferably from 0.03 to 1.5, most preferably from 0.05 to 1.5 percent by weight based upon the weight of the epoxy-containing compound.

The advancement reaction can be conducted at atmospheric, superatmospheric or subatmospheric pressures at temperatures of from 20°C to 260°C, preferably from 80°C to 200°C, more preferably from 100°C to 150°C. The time required to complete the advancement reaction depends upon the temperature employed. Higher temperatures require shorter periods of time whereas lower temperatures require longer periods of time. Generally, however, times of from 5 minutes to 24 hours, preferably from 30 minutes to 8 hours, more preferably from 30 minutes to 3 hours are suitable.

If desired, the advancement reaction can be conducted in the presence of one or more solvents. Suitable such solvents include, for example, glycol ethers, aliphatic and aromatic hydrocarbons, aliphatic ethers, cyclic ethers, ketones, esters, amides, and combinations thereof. Particularly suitable solvents include, for example, toluene, benzene, xylene, methyl ethyl ketone, methyl isobutyl ketone, diethylene glycol methyl ether, dipropylene glycol methyl ether, N,N-dimethylformamide, N-methylpyrrolidinone, tetrahydrofuran, propylene glycol methyl ether and combinations thereof. The solvents can be employed in amounts of from zero to 80 percent, preferably from 20 percent to 60 percent, more preferably from 30 percent to 50 percent by weight based upon the weight of the reaction mixture.

While the advancement of epoxy resins with active hydrogen-containing compounds having rigid, liquid crystal type structures results in an epoxy resin with improved properties, an epoxy resin with even more improvement can be obtained if the advanced epoxy resin compound is subjected to external fields or induced stresses. These fields or stresses provide the orientation of the epoxy resin advancement adducts due to their anisotropic properties. To achieve this orientation during processing, electric or magnetic fields and shear stresses can be applied. The preferred method for orientation is through the application of shear stress. In addition to orientation by electric and magnetic fields, polymeric mesophases can be oriented by low shear rates ($0.1\text{--}90\text{ sec}^{-1}$) which are induced by flow through dies, orifices, and mold gates. For the potential mesomorphic systems based on the advancement of epoxy resins with rigid, liquid crystal type structures, shear orientation can be induced by processing methods such as injection molding, extrusion, filament winding, pultrusion and filming at temperatures from ambient to 200°C , preferably from ambient to 160°C and more preferably from ambient to 120°C . At the time of this processing, an epoxy resin hardener and/or catalyst which would be a part of the final formulated system would set or cure the resin in an oriented, highly ordered state from 1 minute to 1 hour, preferably from 1 minute to 30 minutes, and more preferably from 1 minute to 10 minutes.

The advanced epoxy resins of the present invention can be cured by the conventional means for curing epoxy resins such as by mixing with any suitable curing agent. Suitable such curing agents include, for example, primary and secondary polyamines, carboxylic acids and anhydrides thereof, phenolic hydroxyl-containing compounds, guanidines, urea-aldehyde resins, melamine-aldehyde resins, alkoxyated urea-aldehyde resins, alkoxyated melamine-aldehyde resins, aliphatic, cycloaliphatic and aromatic amines and combinations thereof. Particularly suitable curing agents include, for example, ethylenediamine, diethylenetriamine, triethylenetetramine, tetraethylenepentamine, sulfanilamide, bis(hydroxyphenyl)methane, methylenedianiline, maleic anhydride, chlorendic anhydride, diaminocyclohexane, isophoronediamine, methylbicyclo(2.2.1)heptene-2,3-dicarboxylic anhydride, phenol-formaldehyde novolac resin, cresol-formaldehyde novolac resin, methylolated urea-formaldehyde resin, methylolated melamine-formaldehyde novolac resin, diethyltoluenediamine, diaminodiphenyl sulfone and combinations thereof. The curing agents are employed in an amount which will effectively cure the composition containing the mesogenic moieties. These amounts will depend upon the particular advanced epoxy resin and curing agent employed; however, suitable amounts include, for example, from 0.95:1 to 1.2:1, more suitably from 0.95:1 to 1.15:1, most suitably from 1:1 to 1.15:1 equivalents of curing agent per epoxide equivalent for those curing agents which cure by reacting with the epoxy group of the epoxy resin. The Handbook of Epoxy Resins by Lee and Neville, McGraw-Hill, 1967, contains various discussions concerning the curing of epoxy resins as well as compilation of suitable curing agents.

The advanced epoxy resin compound of the present invention can be blended with other materials such as solvents or diluents, fillers, pigments, dyes, flow modifiers, thickeners, reinforcing agents, mold release agents, wetting agents, stabilizers, fire retardant agents, surfactants and combinations thereof.

These additives are added in functionally equivalent amounts, e.g., the pigments and/or dyes are added in quantities which will provide the composition with the desired color; however, they are suitably employed in amounts of from zero to 20, more suitably from 0.5 to 5, most suitably from 0.5 to 3 percent by weight based upon the weight of the total blended composition.

Solvents or diluents which can be employed herein include, for example, hydrocarbons, ketones, glycol ethers, aliphatic ethers, cyclic ethers, esters, amides and combinations thereof. Particularly suitable solvents or diluents include, for example, toluene, benzene, xylene, methyl ethyl ketone, methyl isobutyl ketone, diethylene glycol methyl ether, dipropylene glycol methyl ether, dimethylformamide, N-methylpyrrolidinone, tetrahydrofuran, propylene glycol methyl ether and combinations thereof.

The modifiers such as thickeners and flow modifiers can be suitably employed in amounts of from zero to 10, more suitably from 0.5 to 6, most suitably from 0.5 to 4 percent by weight based upon the weight of the total composition.

Reinforcing materials which can be employed herein include, for example, natural and synthetic fibers in the form of mats, woven fabric, monofilament, multifilament, unidirectional fiber, rovings, random fibers or filaments, inorganic fillers or whiskers and hollow spheres. Suitable reinforcing materials which can be employed herein include, for example, glass, ceramics, nylon, rayon, cotton, aramid, graphite, polyalkylene terephthalate, polyethylene, polypropylene polyesters and combinations thereof.

Suitable fillers which can be employed herein include, for example, inorganic oxides, ceramic microspheres, plastic microspheres, inorganic whiskers, glass microspheres, CaCO_3 and combinations thereof.

The fillers can be employed in amounts suitably from zero to 95, more suitably from 10 to 80, most suitably from 40 to 60 percent by weight based upon the weight of the total composition.

The following examples are illustrative of the present invention.

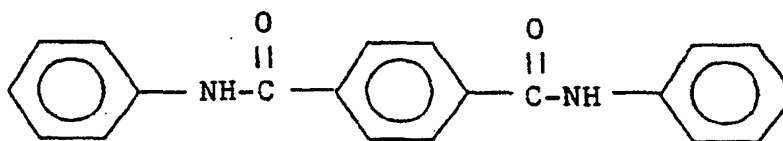
Synthesis of N,N'-Diphenylterephthalate

N,N'-diphenylterephthalamide (Structure I) was synthesized from aniline and terephthaloyl chloride using acetone as a reaction media and an aqueous solution of NaOH to neutralize the HCl generated.

5 Terephthaloyl chloride (87.2 grams, 0.43 moles) was first added to a stirred, 2-liter reaction flask containing one liter of acetone. After the terephthaloyl chloride has dissolved, aniline (80.0 grams, 0.86 moles) was added dropwise using an addition funnel over a 30 minute period. During the course of this addition, a white precipitate was formed. A 200 ml aqueous solution of NaOH (34.3 grams, 0.86 moles) was then added, also using an addition funnel, over a 30 minute period. The reaction temperature during both the aniline and

10 NaOH addition was maintained below 35 °C by blowing air onto the outside of the reaction flask. After the NaOH addition, the reaction mixture was stirred for three hours and then vacuum filtered (filtrate pH = 7). The filter cake obtained was washed with 400 ml of acetone and then added to one liter of deionized water, stirred for 15 minutes and filtered. This filter cake was washed with 500 ml of acetone and then dried in a 105 °C vacuum oven to a constant weight. This final product (114.4 grams, yield = 84.2 percent) exhibited

15 a sharp melting endotherm by differential scanning calorimetry (DSC) at 346 °C. Also Fourier transform infrared (FTIR) analysis showed the following absorbances which are indicative of the structure for the final product: 3329 cm⁻¹ (N-H stretch), 1528 cm⁻¹ (Amide II band) and 1650 cm⁻¹ (Amide I band).



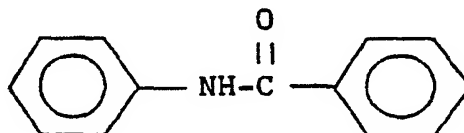
Structure I

Synthesis of N-Phenylbenzamide

N-phenylbenzamide (Structure II) was synthesized from aniline and benzoyl chloride using acetone as a reaction media and an aqueous solution of NaOH to neutralize the HCl generated. A 20 ml aqueous solution of NaOH (2.57 grams, 0.064 moles) and 6.0 grams (0.064 moles) of aniline were first added to a stirred 250

35 ml flask containing 75 ml of acetone. Benzoyl chloride (9.06 grams, 0.064 moles) was next added over a five minute period. This reaction mixture which now contains a white precipitate was stirred for 1.5 hours and then diluted with 50 ml of deionized water. This solution was next filtered and the filter cake obtained was sequentially washed with 50 ml of deionized water (two times) followed by 50 ml of methanol. The washed filter cake was recovered and analysis of this product showed a sharp melting endotherm at 169 °C.

40



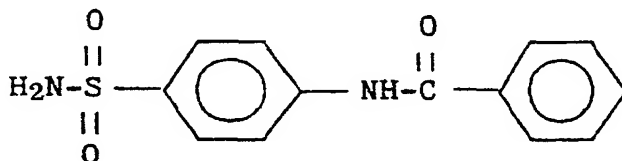
Structure II

Synthesis of 4'-Sulfonamido-N-Phenylbenzamide

4'-sulfonamido-N-phenylbenzamide (Structure III) was synthesized from sulfanilamide and benzoyl chloride using acetone as a reaction media and an aqueous solution of NaOH to neutralize the HCl generated. Sulfanilamide (100 grams, 0.58 moles) was first added to a stirred, two liter reaction flask containing one liter of acetone. After the sulfanilamide had dissolved, benzoyl chloride (82 grams, 0.58

55

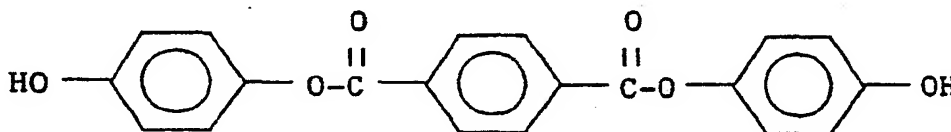
moles) was added dropwise using an addition funnel over a 20 minute period. During the course of this addition, a white precipitate was formed. A 200 ml aqueous solution of NaOH (23.2 grams, 0.58 moles) was then added, also using an addition funnel, over a 30 minute period. The reaction temperature during this addition and that of the benzoyl chloride was maintained below 30 °C by blowing air onto the outside of the reaction flask. After the NaOH addition, the reaction mixture was stirred for two hours and then vacuum filtered. The filter cake obtained was washed with one liter of deionized water. These solids were next added to 300 ml of acetone, stirred for 15 minutes, and then vacuum filtered. This filter cake was washed with 500 ml of deionized water followed by 300 ml of acetone and then dried to a constant weight in a 100 °C vacuum oven. The final dried product (145 grams, yield = 90.2 percent) exhibited a sharp melting endotherm by DSC at 297 °C. Also FTIR analysis of this material showed the following absorbances which are indicative of the structure for the final product: 3350 cm⁻¹ (N-H stretch), 1520 cm⁻¹ (Amide II band), 1650 cm⁻¹ (Amide I band) and 3267 cm⁻¹/3291 cm⁻¹ (N-H stretch of sulfonamide).



Structure III

Synthesis of Bis(4-Hydroxyphenyl)Terephthalate

Bis(4-hydroxyphenyl)terephthalate (Structure IV) was synthesized by the reaction of hydroquinone with terephthaloyl chloride through the addition of NaOH. The hydroquinone (162.7 grams, 1.48 moles) and terephthaloyl chloride (150.0 grams, 0.74 moles) were first dissolved in 750 ml of tetrahydrofuran in a two liter round bottom flask which was stirred. The NaOH (59.1 grams, 1.48 moles) in 300 ml of deionized water was next added dropwise using an addition funnel over a one hour period and the mixture was then allowed to stir an additional two hours at ambient temperature (pH = 6). The precipitate obtained was collected by vacuum filtration and then added to 500 ml of deionized water and 1000 ml of methanol. This solution was stirred for one hour at 40-55 °C and then filtered. The solids collected here were added to 1100 ml of anhydrous methanol and stirred at 40-55 °C for one hour followed by a third filtration. The resultant solids were then dried in a 80 °C vacuum oven. This final product (111.9 grams, yield = 43.3 percent) exhibited a sharp melting endotherm by DSC at 415 °C.

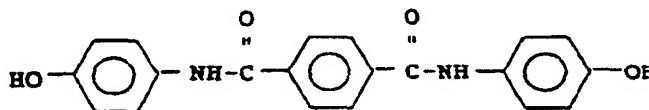


Structure IV

Synthesis of N,N'-Bis(4-Hydroxyphenyl)Terephthalamide

N,N'-bis(4-hydroxyphenyl)terephthalamide (Structure V) was synthesized by the reaction of aminophenol with terephthaloyl chloride in the presence of NaHCO₃. Two moles of aminophenol (218.3 grams) and 2 moles of NaHCO₃ (168 grams) were first added to a stirred, round bottom flask containing 750 milliliters of acetone and 750 milliliters of deionized water. Two moles of terephthaloyl chloride (203.0 grams) were then added over a two hour period. During the course of the terephthaloyl chloride addition, 200 additional milliliters of acetone were also added to wash down the frothy suspension caused by CO₂

evolution. After the terephthaloyl chloride addition, the precipitate produced from the reaction was collected by vacuum filtration. For purification of the precipitate obtained, one half of this material was placed in a stirred, round bottom flask containing 900 milliliters of methanol and 450 milliliters of deionized water. This suspension was heated to 56 °C and then vacuum filtered. For the solids recovered from this filtration, the washing step with methanol and deionized water was repeated. The filtrant solids obtained from the second hot filtration were then added to a stirred, round bottom flask containing 1250 milliliters of methanol. This suspension was vacuum filtered after 30 minutes to recover the solids for drying. After following the preceding washing procedure for the second half of the reaction precipitate, the total solids recovered were dried at 80 °C under vacuum for 5 hours. The weight of the dried solids was 325.6 grams, 93.4 percent of the theoretical yield to N,N'-bis(4-hydroxyphenyl)terephthalamide. The melting point of this final product was 405 °C as determined by differential scanning calorimetry.



Structure V

Comparative Experiment A

Cured Low Molecular Weight Epoxy Resin

A diglycidyl ether of bisphenol A, 149.1 grams having 24.6 percent epoxide and a 174.8 epoxide equivalent weight (EEW) was heated to 120 °C. A conventional epoxy resin hardener, diethyltoluenediamine (39.51 grams, 1.04 times the stoichiometric amount of epoxide) was then added. After mixing, this resin system was poured into a mold (dimensions = 8 inches x 8 inches x 1/8 inch; 203.2 mm x 203.2 mm x 3.175 mm) heated to 120 °C in a convection oven. The temperature of the oven was then increased to 180 °C. After two hours at 180 °C, the oven was cooled to room temperature. Following cool down, a void free, neat resin casting was obtained from the mold. Glass transition temperature and mechanical properties for this casting were then determined. These results are reported in Table I and again in Table II for comparison purposes.

Comparative Experiment B

Cured Advanced Epoxy Resin with Bisphenol A -Control

A diglycidyl ether of bisphenol A, 150 grams (1 equiv.) having a percent epoxide of 16.3 and an EEW of 263.8 advanced with 25.1 grams (0.25 equiv.) of bisphenol A was heated to 120 °C. A stoichiometric amount (29.25 grams) of diethyltoluenediamine was then added. After mixing, this resin system was poured into a mold (dimensions = 8 inches x 8 inches x 1/8 inch; 203.2 mm x 203.2 mm x 3.175 mm) heated to 120 °C in a convection oven. The temperature of the oven was then raised to 180 °C. After five hours at 180 °C, the oven was cooled to room temperature. Following cool down, a void free, neat resin casting was obtained from the mold. Glass transition temperature and mechanical properties for this casting were then determined. These results are reported in Table III.

Example 1 Advancement of Diglycidyl Ether of Bisphenol A Resin with N,N'-Diphenylterephthalamide

N,N'-diphenylterephthalamide (9.00 grams) was added to a stirred, 500 ml reaction flask containing 300.1 grams of a diglycidyl ether of bisphenol A having an EEW of 177.7. A sample of this material when viewed under an optical microscope (70 X magnification) using a cross polarized light source shows

dispersed crystallites of the N,N'-diphenylterephthalamide in the epoxy resin. This mixture was then heated to 248 °C and held at this temperature for 1.5 hours. After cooling to room temperature, optical microscopy shows birefringent domains in the epoxy resin. The percent epoxide of this advanced resin was 22.7 based on titrametric determination. This percent epoxide corresponds to the expected value obtained by calculations based on the complete reaction of the amide hydrogens of N,N'-diphenylterephthalamide.

Example 2 Cure of Diglycidyl Ether of Bisphenol A Epoxy Resin Advanced with N,N'-Diphenylterephthalamide

Part of the advanced epoxy resin from Example 1 (147.0 grams) was heated to 120 °C. Diethyltoluenediamine (35.81 grams, 1.04 times the stoichiometric amount of epoxide) was then added. After mixing, this resin system was poured into a mold (dimensions = 8 inches x 8 inches x 1/8 inch; 203.2 mm x 203.2 mm x 3.175 mm) heated to 120 °C in a convection oven. The temperature of the oven was then raised to 180 °C. After two hours at 180 °C, the oven was cooled to room temperature and a void free, neat resin casting was obtained from the mold. When this casting was viewed under an optical microscope (70 X magnification) using a cross polarized light source, rodlike, birefringent segments were observed which were dispersed throughout the polymer. Glass transition temperature and mechanical properties were then determined for this casting and these results are reported in Table I. Compared to the diglycidyl ether of bisphenol A epoxy resin control of Example 1, the cured resin shows improvements in tensile and flexural modulus of 4.7 and 5.9 percent, respectively.

Example 3 Cure of Diglycidyl Ether of Bisphenol A Epoxy Resin Advanced with N,N'-Diphenylterephthalamide (Orientation of Birefringent Segments with the Application of an Electric Field)

A diglycidyl ether of bisphenol A epoxy resin advanced with N,N'-diphenylterephthalamide (143.3 grams, 22.5 percent epoxide) was first heated to 160 °C. Diethyltoluenediamine (34.86 grams, 1.04 times the stoichiometric amount of epoxide) was then added. After mixing, this resin system was poured into an aluminum mold (dimensions = 8 inches x 8 inches x 1/8 inch; 203.2 mm x 203.2 mm x 3.175 mm) heated to 120 °C in a convection oven. The mold consisted of two aluminum plates spaced apart by an 1/8 inch (3.175 mm) silicone rubber gasket which formed the mold cavity. Prior to pouring of the resin system into the mold, a 300 volt charge was established between the two aluminum plates using a direct current power supply. After one hour at 120 °C, the oven temperature was raised to 180 °C and held at this temperature for two hours. Following cool down from 180 °C to room temperature, a void free, neat resin casting was obtained from the mold. When this casting was viewed under an optical microscope using a cross polarized light source, orientation of the birefringent segments produced through the advancement of diglycidyl ether of bisphenol A epoxy resin with N,N'-diphenylterephthalamide was observed. This orientation was normal to the electric field application. From this neat resin casting, glass transition temperature and mechanical properties were determined. These results are reported in Table I. Compared to the diglycidyl ether of bisphenol A epoxy resin control of Example 1, improvements in tensile and flexural modulus of 13.5 and 6.4 percent were observed, respectively.

Example 4 Advancement of Diglycidyl Ether of Bisphenol A Epoxy Resin with N,N'-Diphenylterephthalamide and N-Phenylbenzamide

N,N'-diphenylterephthalamide (9.0 grams) and N-phenylbenzamide (3.0 grams) were added to a stirred, 500 ml reaction flask containing 300.0 grams of diglycidyl ether of bisphenol A epoxy resin (24.5 percent epoxide). This mixture was then heated to 248 °C and held at this temperature for approximately 1.5 hours. The measured percent epoxide of this advanced resin was 22.4 which compared to a calculated percent epoxide of 22.3 based on complete reaction of the amide hydrogens of N,N'-diphenylterephthalamide and N-phenylbenzamide. After 1.5 hours at 248 °C, the resin was cooled quickly to room temperature by pouring into a large aluminum pan. When this resin was viewed under an optical microscope (70 X magnification) using a cross polarized light source, dispersed birefringent areas were observed.

Example 5 Cure of Diglycidyl Ether of Bisphenol A Epoxy Resin Advanced with N,N'-Diphenyltereph-

thalamide and N-Phenylbenzamide

Part of the advanced diglycidyl ether of bisphenol A epoxy resin of Example 4 (150.1 grams) was heated to 120° C. Diethyltoluenediamine (36.20 grams, 1.04 times the stoichiometric amount of epoxide) was then added. After mixing, this resin system was poured into an aluminum mold heated to 120° C in a convection oven. The configuration of this aluminum mold was the same as in Example 3 and after pouring the resin into this mold, a 16 volt charge was established between the aluminum plates using a direct current power supply. After two hours at 120° C, the oven was raised to 180° C and held for two hours at this temperature. Following cool down of the oven from 180° C to room temperature, a void free, neat resin casting was obtained from the mold. When this casting was viewed under an optical microscope using a cross polarized light source, dispersed birefringent particles were observed in the polymer. The glass transition temperature and mechanical properties for this casting were then determined and these results are reported in Table I. Compared to the diglycidyl ether of bisphenol A epoxy resin control of Example 1, the cured resin shows an improvement in tensile and flexural modulus of 9.7 and 15.2 percent, respectively.

Example 6 Advancement of Diglycidyl Ether of Bisphenol A Epoxy Resin with 4'-Sulfonamido-N-Phenylbenzamide

Thirty (30) grams of 4'-sulfonamido-N-phenylbenzamide were added to a stirred, 500 ml reaction flask containing 300.7 grams of diglycidyl ether of bisphenol A epoxy resin (24.4 percent epoxide). A sample of this mixture when viewed under an optical microscope (70 X magnification) using a cross polarized light source showed dispersed crystallites of 4'-sulfonamido-N-phenylbenzamide in the epoxy resin. This mixture was then heated. At 218° C, the mixture became clear and a small exotherm was observed. Following the exotherm, which peaked at 235° C, the resin was cooled to room temperature. At room temperature, no solids were observed. When viewed under the optical microscope (70 x magnification), small birefringent areas were observed in the resin. The measured percent epoxide of this resin was 18.5 which compared to a calculated percent epoxide of 18.0 based on the complete reaction of all the active hydrogens of the 4'-sulfonamido-N-phenylbenzamide.

Example 7 Advancement of Diglycidyl Ether of Bisphenol A Epoxy Resin with 4'-Sulfonamido-N-Phenylbenzamide

Forty-five (45) grams of 4'-sulfonamido-N-phenylbenzamide were added to a stirred, 500 ml reaction flask containing diglycidyl ether of bisphenol A epoxy resin (299.8 grams, 24.8 percent epoxide). This mixture was then heated. At 220° C the mixture started to clear and a small exotherm was observed. Following the exotherm, which peaked at 248° C, the resin was cooled to room temperature. At room temperature, no solids were observed in the resin. When viewed under an optical microscope (70 X magnification) using a cross polarized light source, small birefringent areas could be seen. The titrated percent epoxide of this advanced resin was 16.3 which compared to a calculated percent epoxide of 15.5 based on the complete reaction of all the active hydrogens of the 4-sulfonamido-N-phenylbenzamide.

Example 8 Cure of Diglycidyl Ether of Bisphenol A Epoxy Resin Advanced with 4'-Sulfonamido-N-Phenylbenzamide

Part of the resin obtained from Example 6 (145.8 grams) was heated to 120° C. Diethyltoluenediamine (29.1 grams, 1.04 times the stoichiometric amount of epoxide) was then added. After mixing, this resin system was poured into a mold (dimensions = 8 inches x 8 inches x 1/8 inch; 203.2 mm x 203.2 mm x 3.175 mm) heated to 120° C in a convection oven. The temperature of the oven was then raised to 180° C and held at this temperature for two hours. Following cool down of the oven from 180° C to room temperature, a void free, neat resin casting was obtained from the mold. The glass transition temperature and mechanical properties of this casting were then determined and these results are reported in Table II. Compared to the diglycidyl ether of bisphenol A epoxy resin control of Example 1, this cured resin showed improvements in tensile and flexural modulus of 10.9 and 9.3 percent, respectively.

Example 9 Cure of Diglycidyl Ether of Bisphenol A Epoxy Resin Advanced with 4'-Sulfonamido-N-Phenylbenzamide (Orientation of Birefringent Segments with the Application of an Electric Field)

Part of the resin obtained from Example 6 (174.8 grams) was heated to 120° C. Diethyltoluenediamine (30.82 grams, 1.04 times the stoichiometric amount of epoxide) was then added. After mixing, this resin system was poured into an aluminum mold heated to 120° C in a convection oven. The configuration of this mold was the same as in Example 3 and after pouring of the resin, a 16 volt charge was established between the aluminum plates using a direct current power supply. After two hours at 120° C, the oven temperature was raised to 180° C. After two hours at 180° C, the oven was cooled to room temperature and a void free, neat resin casting was obtained from the mold. When this casting was viewed under an optical microscope (120 X magnification) using a cross polarized light source, birefringent fibers were observed in the polymer. The glass transition temperature and mechanical properties for this casting were then determined and these results are reported in Table II. Compared to the diglycidyl ether of bisphenol A epoxy resin control of Example 1, this cured resin shows improvements in tensile and flexural modulus of 13.5 and 11.3 percent, respectively.

Example 10 Cure of Diglycidyl Ether of Bisphenol A Epoxy Resin Advanced with 4'-Sulfonamido-N-Phenylbenzamide

A diglycidyl ether of Bisphenol A epoxy resin which had been advanced with 4'-sulfonamido-N-phenylbenzamide in the same manner as Example 6 was heated to 120° C. The weight of this advanced resin, which contained 18.7 percent epoxide, was 169.9 grams. Diethyltoluenediamine (34.21 grams, 1.04 times the stoichiometric amount of epoxide) was then added. After mixing, this resin system was poured into an aluminum mold heated to 120° C in a convection oven. The configuration of this mold was the same as in Example 3. After pouring the resin into the mold, a 50 volt charge was established between the aluminum plates using a direct current power supply. After one hour at 120° C, the oven temperature was raised to 180° C. After two hours at 180° C, the oven was cooled to room temperature and a void free, neat resin casting was obtained from the mold. The glass transition temperature and mechanical properties for this casting were then determined and these results are reported in Table II. Compared to the diglycidyl ether of bisphenol A epoxy resin control of Example 1, this cured resin shows improvements in tensile and flexural modulus of 16.4 and 12.4 percent, respectively. In addition to achieving these higher moduli, the strain to failure for this polymer is equal to that of the control.

Example 11 Cure of Diglycidyl Ether of Bisphenol A Epoxy Resin Advanced with 4'-Sulfonamido-N-Phenylbenzamide

Part of the resin from Example 7 (174.8 grams) was heated to 120° C. Diethyltoluenediamine (30.82 grams, 1.04 times the stoichiometric amount of epoxide) was then added. After mixing, this resin system was poured into an aluminum mold heated to 120° C in a convection oven. The configuration of this mold was the same as in Example 3. After pouring the resin into the mold, a 300 volt charge was established between the aluminum plates using a direct current power supply. After one hour at 120° C, the temperature of the oven containing the mold was raised to 180° C and held for two hours at this temperature. Following cool down of the oven from 180° C to room temperature, a void free, neat resin casting was obtained from the mold. The glass transition temperature and mechanical properties for this casting were then determined and these results are reported in Table III. Compared to diglycidyl ether of bisphenol A epoxy resin advanced with bisphenol A (Comparative Experiment B), this cured resin shows improvements in tensile and flexural modulus of 17.9 and 18.5 percent, respectively.

Example 12 Advancement of Diglycidyl Ether of Bisphenol A Epoxy Resin with Bis(4-Hydroxyphenyl)Terephthalate Followed by Cure with Diethyltoluenediamine

Bis(4-hydroxyphenyl)terephthalate (9.01 grams) was mixed into 50.0 grams of diglycidyl ether of bisphenol A epoxy resin containing 2500 ppm tetrabutylphosphonium acetate*acetic acid complex (70 weight percent in methanol) catalyst. This mixture, which was contained in a stirred flask, was then heated to 140° C. Prior to heating, differential scanning calorimetry analysis of this mixture showed a reaction

exotherm beginning at 120°C (peak temperature = 170°C, $\Delta H = -64$ joules/gram). After three hours at 140°C, the temperature was reduced to 120°C. The resin at this point contained dispersed crystalline segments as observed by optical microscopy (70 X magnification) using a cross polarized light source. Diethyltoluenediamine (10.82 grams, 1.04 times the calculated stoichiometric amount of epoxide remaining after complete advancement) was then added. After mixing, this resin system was poured into an aluminum mold heated to 120°C in a convection oven. The mold used was of the same configuration as used in Example 3. After pouring the resin into the mold, a 7 volt charge was applied between the two aluminum plates using a direct current power supply. The oven temperature was then raised to 180°C where it was maintained for four hours. Following cool down of the oven from 180°C to room temperature, a void free, translucent casting was obtained from the mold. The glass transition temperature of this polymer was 131°C as determined by differential scanning calorimetry which also indicates no additional reactivity to 300°C. The flexural strength and modulus for the casting were 17,680 psi and 463 ksi, respectively. Compared to diglycidyl ether of bisphenol A epoxy resin advanced with bisphenol A (Comparative Experiment B), the flexural modulus obtained represents a 16.9 percent improvement.

Example 13 Advancement of the Diglycidyl Ether of Bisphenol A with N,N'-Bis(4-hydroxyphenyl)-terephthalamide and Preparation of a Compression Molded Phenoxo Resin Casting

N,N'-Bis(4-hydroxyphenyl)terephthalamide (30.7 grams; 0.176 hydroxyl equivalent) was blended into an equivalent amount of the diglycidyl ether of bisphenol A (30.0 grams; EEW = 170.3; 0.176 epoxide equivalent) which contained 4500 ppm tetrabutylphosphonium acetate*acetic acid complex (70 weight percent in methanol) catalyst. This blend was then placed in a 130°C convection oven where it was stirred approximately every 5 minutes. Differential scanning calorimetry analysis of a sample of this blend showed a reaction exotherm ($\Delta H = -120$ joules/gram) with an onset temperature of 160°C. The peak temperature of this exotherm was 240°C. After 30 minutes at 130°C, the oven temperature was increased to 180°C over a 1.5 hour period during which time periodic stirring was continued. After 30 minutes at 180°C, the thickened resin was removed from the oven and cooled to approximately 0°C. The cooled, solid resin was then ground to a fine powder. For the preparation of a neat resin casting, part of this powder (54 grams) was placed in a compression mold (dimensions = 4 inches x 4 inches x 0.125 inch; 101.6 mm x 101.6 mm x 3.18 mm). This compression mold was next transferred to a mechanical press heated to 220°C. In the press, pressure (1680 psi) was applied to the mold during the first 5 minutes. After 1.5 hours at 220°C, the temperature of the press was increased to 240°C where it was maintained for 1.5 hours before cooling to room temperature. At room temperature, an opaque, neat resin casting was obtained from the mold. The melting point of this polymer was 378°C as determined by differential scanning calorimetry. The flexural strength and modulus of the neat resin casting obtained were 8440 psi and 713 ksi, respectively.

TABLE I

DIGLYCIDYL ETHER OF BISPHENOL A EPOXY RESIN ADVANCED WITH N,N'-DIPHENYLTEREPHTHALAMIDE AND N-PHENYLBENZAMIDE - PROPERTIES FOR NEAT RESIN CURED WITH DIETHYLTOLUENEDIAMINE				
EXAMPLE NO.	Comp. Expt. A	2	3	5
Wt. % N,N'-Diphenylterephthalamide Used in Advancement Reaction	0	2.9	2.9	2.9
Wt. % N-Phenylbenzamide Used in Advancement Reaction	0	0	0	0.96
% Epoxide of Resin	24.6	22.7	22.5	22.4
Electric Field Applied During Cure, volts D.C.	0	0	300	16
Glass Transition Temperature, °C	179	173	169	165
Tensile Strength, psi	10,560	9,920	7,605	9,915
Tensile Modulus, ksi	340	356	386	373
Strain to Failure, %	5.3	3.8	2.4	3.9
Flexural Strength, psi	19,320	13,060	13,360	18,420
Flexural Modulus, ksi	388	411	413	447

TABLE II

DIGLYCIDYL ETHER OF BISPHENOL A EPOXY RESIN ADVANCED WITH 4'-SULFONAMIDO-N-PHENYLBENZAMIDE - PROPERTIES FOR NEAT RESIN CURED WITH DIETHYLTOLUENEDIAMINE				
EXAMPLE NO.	Comp. Expt. A	8	9	10
Wt. % 4'-Sulfonamido-N-Phenylbenzamide Used in Advancement Reaction	0	9.1	9.1	9.1
% Epoxide of Resin	24.6	18.5	18.5	18.7
Electric Field Applied During Cure, volts D.C.	0	0	16	50
Glass Transition Temperature, °C	179	168	166	167
Tensile Strength, psi	10,560	8,810	11,120	11,740
Tensile Modulus, ksi	340	377	386	416
Strain to Failure, %	5.3	3.1	4.8	5.2
Flexural Strength, psi	19,320	20,970	20,180	20,850
Flexural Modulus, ksi	388	424	432	436

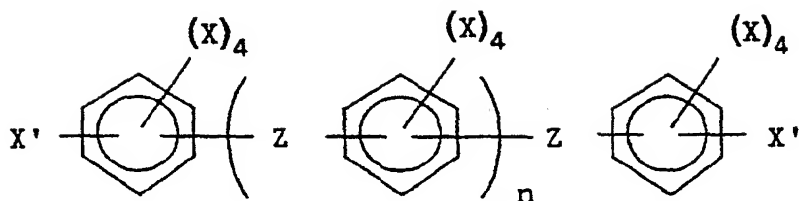
TABLE III

DIGLYCIDYL ETHER OF BISPHENOL A EPOXY RESIN ADVANCED WITH 4'-SULFONAMIDO-N-PHENYLBENZAMIDE - PROPERTIES FOR NEAT RESIN CURED WITH DIETHYLTOLUENEDIAMINE		
EXAMPLE NO.	1	11
Wt. 4'-Sulfonamido-N-Phenylbenzamide Used in Advancement Reaction	0	18.4
Wt. % Bisphenol A Used in Advancement Reaction	14.3	0
% Epoxide of Resin	16.3	16.3
Electric Field Applied During Cure, volts D.C.	0	300
Glass Transition Temperature, °C	150	156
Tensile Strength, psi	11,100	11,510
Tensile Modulus, ksi	351	414
Strain to Failure, %	7.3	4.3
Flexural Strength, psi	19,460	22,360
Flexural Modulus, ksi	396	460

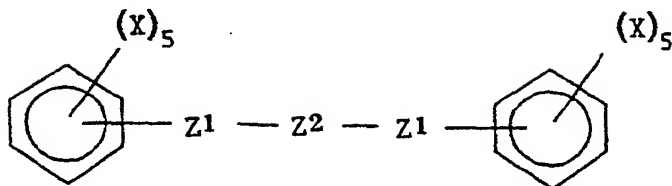
Claims

1. An advanced epoxy resin compound prepared by reacting (A) at least one epoxy resin having an average of more than one vicinal epoxy group per molecule with (B) at least one compound having an average of more than one active hydrogen atom per molecule represented by the following formulas I, II or III:

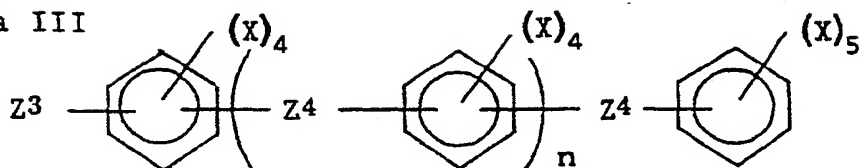
Formula I



Formula II

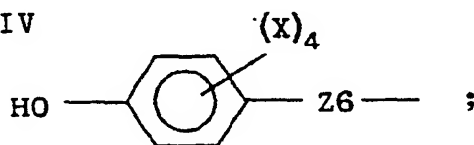


Formula III

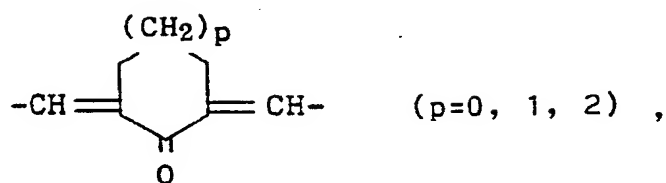
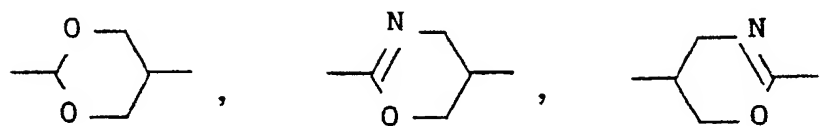
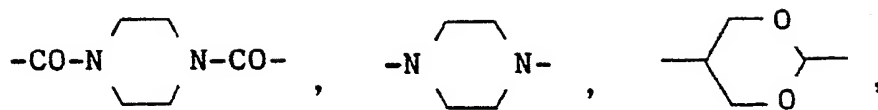
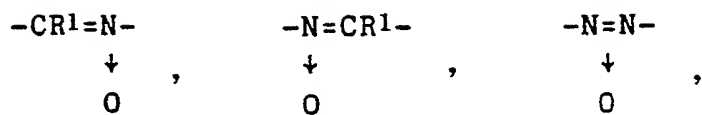


wherein each X is independently hydrogen, a hydrocarbyl or hydrocarbyloxy group having from 1 to 12, preferably from 1 to 6, most preferably from 1 to 4 carbon atoms, a halogen atom, $-\text{NO}_2$ or $-\text{C}\equiv\text{N}$; each X' is independently a hydroxyl group, a carboxylic acid group or the group represented by the following formula IV:

Formula IV



each Z is independently $-\text{CR}^1 = \text{CR}^1-$, $-\text{CR}^1 = \text{CR}^1 - \text{CR}^1 = \text{CR}^1-$, $-\text{CR}^1 = \text{N} = \text{N} = \text{CR}^1-$, $-\text{CR}^1 = \text{CR}^1 - \text{CO} - \text{O} - \text{CH}_2-$, $-\text{CR}^1 = \text{CR}^1 - \text{CO} - \text{O} - \text{CH}_2 - \text{CH}_2-$, $-\text{CH}_2 - \text{O} - \text{CO} - \text{CR}^1 = \text{CR}^1-$, $-\text{CH}_2 - \text{CH}_2 - \text{O} - \text{CO} - \text{CR}^1 = \text{CR}^1-$, $-\text{CR}^1 = \text{CR}^1 - \text{CO} - \text{O}-$, $-\text{O} - \text{CO} - \text{CR}^1 = \text{CR}^1-$, $-\text{N} = \text{N}-$, $-\text{CO} - \text{NH}-$, $-\text{NH} - \text{CO}-$, $-\text{CO} - \text{NH} - \text{NH} - \text{CO}-$, $-\text{C}\equiv\text{C}-$, $-\text{C}\equiv\text{C} - \text{C}\equiv\text{C}-$, $-\text{CO} - \text{S}-$, $-\text{S} - \text{CO}-$, $-\text{CO} - \text{O}-$, $-\text{O} - \text{CO}-$, a direct single bond when $n \geq 1$,



5

10

15

20

25

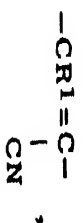
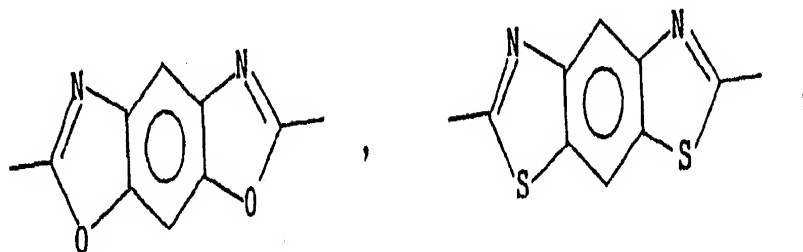
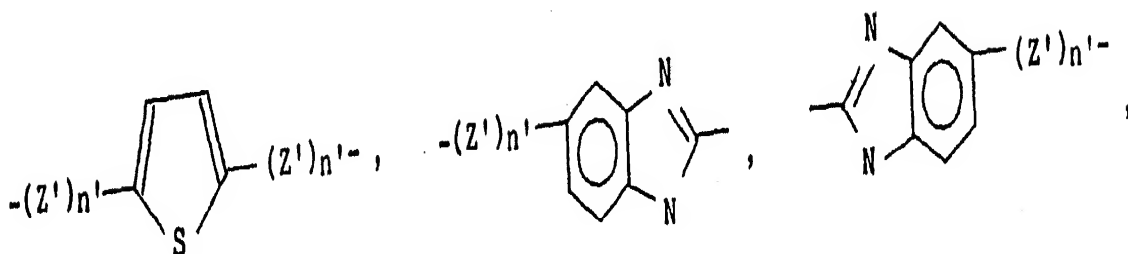
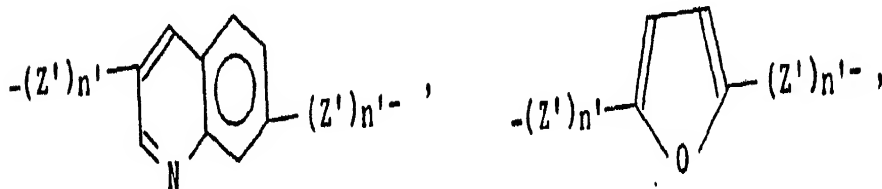
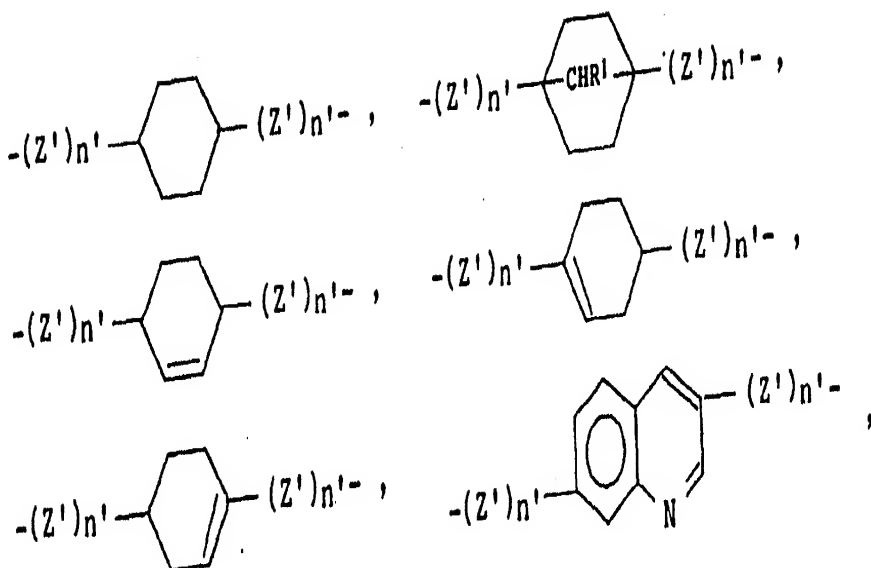
30

35

40

45

50

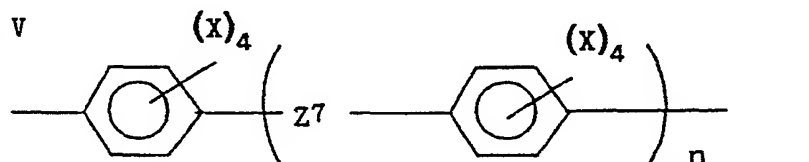


and

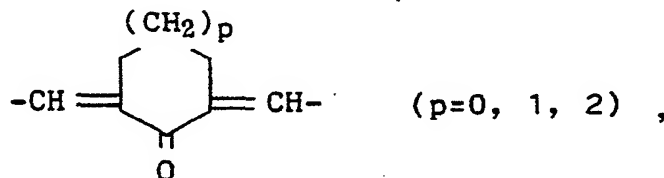
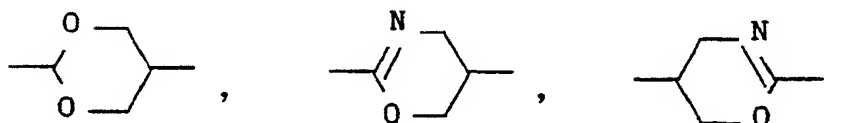
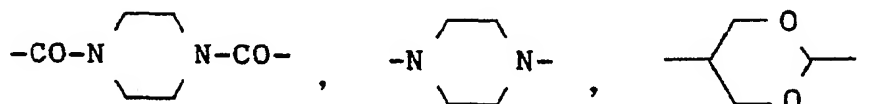
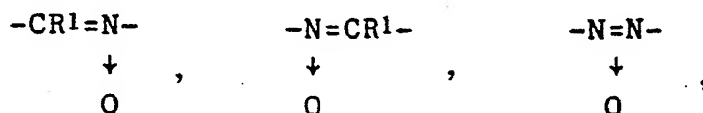


Z' is independently $-O-CO-$, $-CO-O-$, $-NR^1-CO-$, $-CO-NR^1-$; each n' is independently zero or 1; R^1 is independently a hydrogen atom or an alkyl group having from 1 to 4 carbon atoms; Z^6 is a divalent hydrocarbyl group having from 1 to 12, preferably from 1 to 6, more preferably from 1 to 4 carbon atoms; $-SO-$, $-SO_2-$, $-S-$, $-S-S-$, $-O-$, or $-CO-$; each Z^1 is independently $-CO-NH-$, or $-NH-CO-$; Z^2 is a group represented by a cyclic or bicyclic ring system containing from 5 to 12, preferably from 5 to 10, more preferably from 5 to 6 carbon atoms or a group represented by the following formula V:

Formula V



Z^3 is NH_2- , NH_2-SO_2- , NH_2-CO- , or NH_2-Z^5-O- ; each Z^4 is independently $-CR^1=CR^1-$, $-CR^1=CR^1-$, $-CR^1=CR^1-$, $-CR^1=N-N=CR^1-$, $-CR^1=CR^1-CO-O-CH_2-$, $-CR^1=CR^1-CO-O-CH_2-CH_2-$, $-CH_2-O-CO-CR^1=CR^1-$, $-CH_2-CH_2-O-CO-CR^1=CR^1-$, $-CR^1=CR^1-CO-O-$, $-O-CO-CR^1=CR^1-$, $-N=N-$, $-CO-NH-$, $-NH-CO-$, $-CO-NH-$, $-NH-CO-$, $-C\equiv C-$, $-C\equiv C-C\equiv C-$, $-CO-S-$, $-S-CO-$, $-CR^1=N-$, $-N=CR^1-$, $-CO-O-$, $-O-CO-$, a direct single bond,



5

10

15

20

25

30

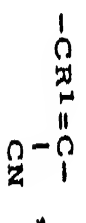
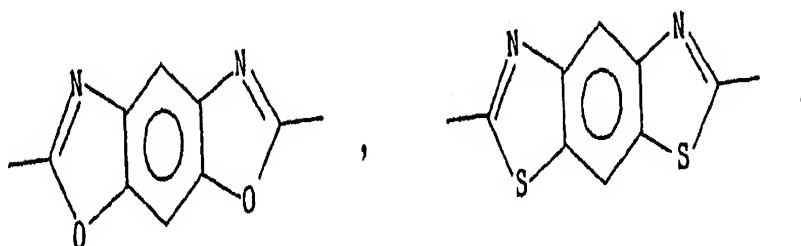
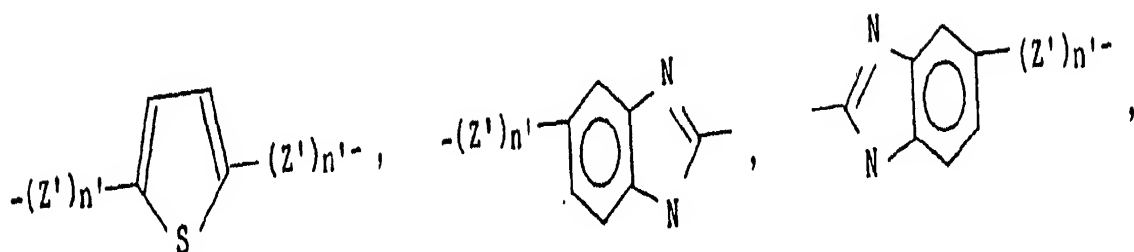
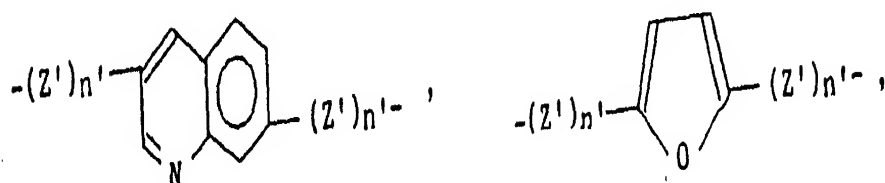
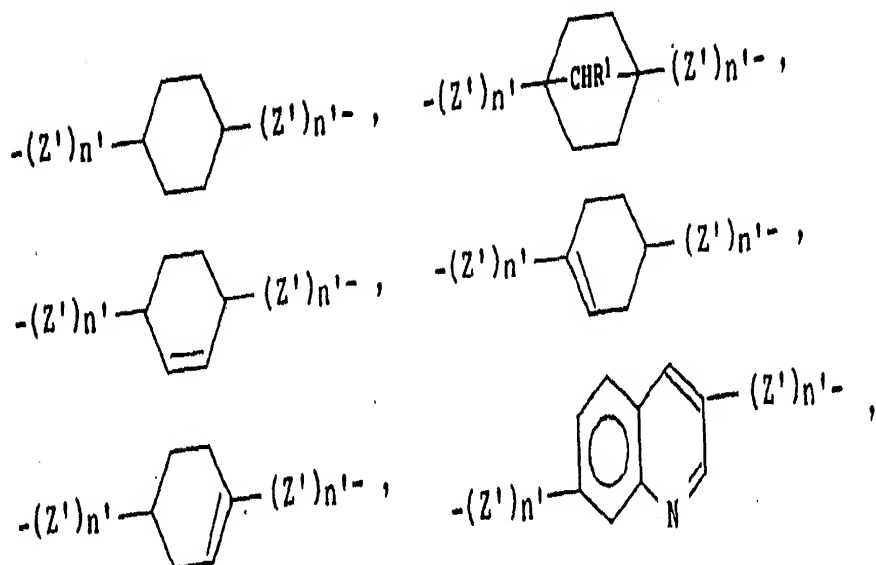
35

40

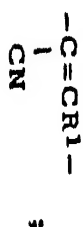
45

50

55



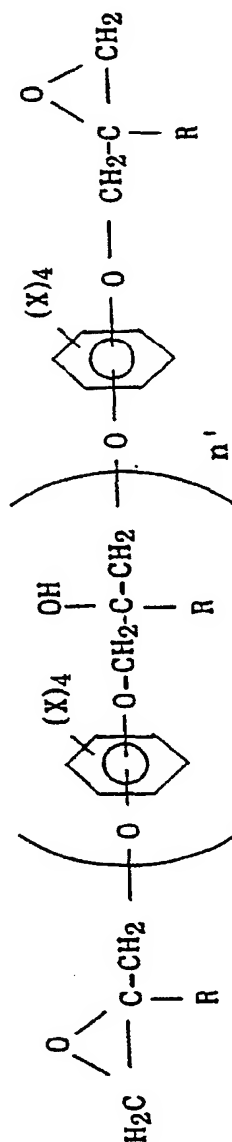
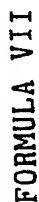
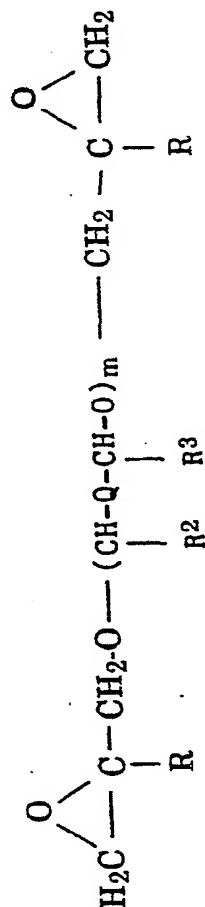
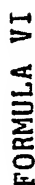
and



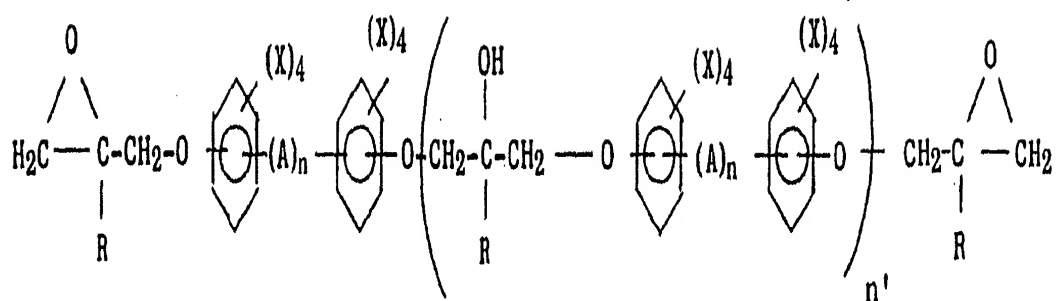
Z⁵ is an alkyl or cycloalkyl group having from 1 to 10, preferably from 1 to 6, more preferably from 1 to 3 carbon atoms; Z⁷ is the same as Z⁴ with the proviso that it can also independently be a divalent hydrocarbonyl group having from 1 to 4, carbon atoms, -SO-, -SO₂-, -S-, -S-S-, -O-, or -CO-; and n has an average value of zero to 6; with the proviso that (a) at least 80 percent of the Z and X' groups are in the para position with respect to each other in formula I and (b) at least 80 percent of the Z³ and Z⁴ groups are in the para position with respect to each other in formula III; and wherein components (A) and (B) are employed in amounts which provide a ratio of active hydrogen atoms per vicinal epoxy group of from 0.01:1 to 1.05:1.

2. An advanced epoxy resin compound of Claim 1 wherein component (B) is 4'-sulfonamido-N-phenylbenzamide or N,N'-bis(4-hydroxyphenyl)terephthalamide.

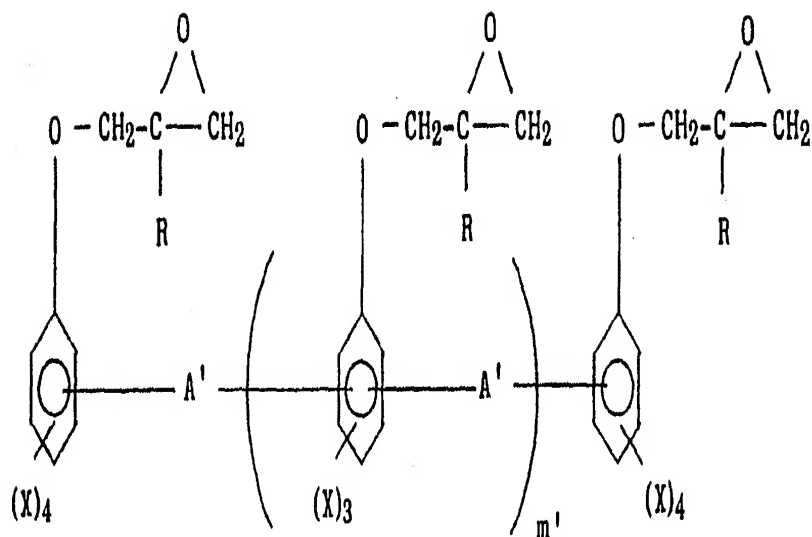
3. An advanced epoxy resin compound of Claim 1 or 2 wherein component (A) is an epoxy resin represented by the following formulas VI, VII, VIII, IX or X:



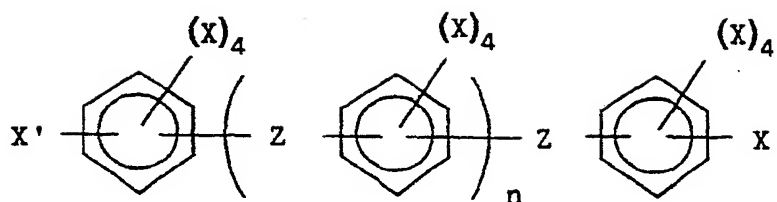
FORMULA VIII



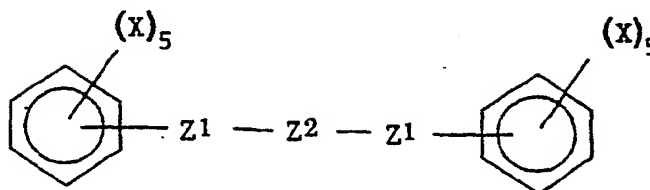
FORMULA IX



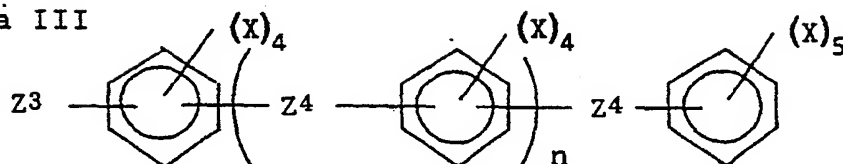
Formula I



Formula II

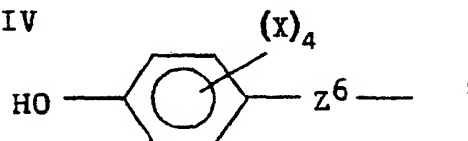


Formula III

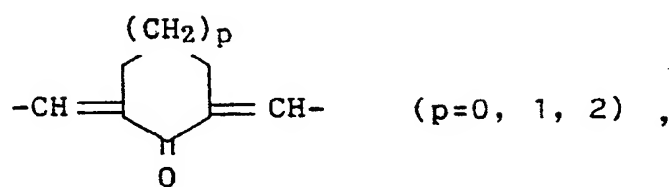
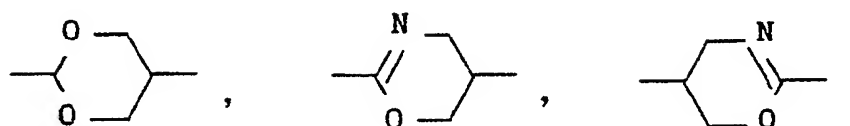
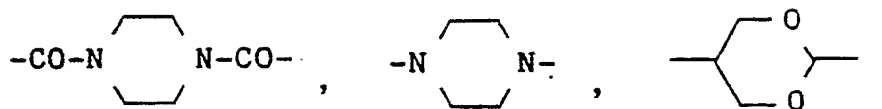
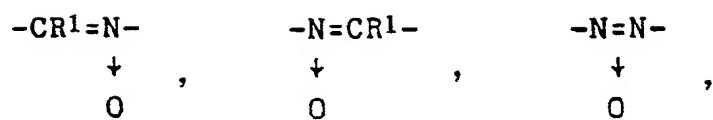


wherein each X is independently hydrogen, a hydrocarbyl or hydrocarbyloxy group having from 1 to 12, preferably from 1 to 6, most preferably from 1 to 4 carbon atoms, a halogen atom, $-\text{NO}_2$ or $-\text{C}\equiv\text{N}$; each X' is independently a hydroxyl group, a carboxylic acid group or the group represented by the following formula IV:

Formula IV



each Z is independently $-\text{CR}'=\text{CR}'$ -, $-\text{CR}'=\text{CR}'-\text{CR}'=\text{CR}'$ -, $-\text{CR}'=\text{N}-\text{N}=\text{CR}'$ -, $-\text{CR}'=\text{CR}'-\text{CO}-\text{O}-\text{CH}_2$ -, $-\text{CR}'=\text{CR}'-\text{CO}-\text{O}-\text{CH}_2-\text{CH}_2$ -, $-\text{CH}_2-\text{O}-\text{CO}-\text{CR}'=\text{CR}'$ -, $-\text{CH}_2-\text{CH}_2-\text{O}-\text{CO}-\text{CR}'=\text{CR}'$ -, $-\text{CR}'=\text{CR}'-\text{CO}-\text{O}$ -, $-\text{O}-\text{CO}-\text{CR}'=\text{CR}'$ -, $-\text{N}=\text{N}$ -, $-\text{CO}-\text{NH}$ -, $-\text{NH}-\text{CO}$ -, $-\text{CO}-\text{NH}-\text{NH}-\text{CO}$ -, $-\text{C}\equiv\text{C}$ -, $-\text{C}\equiv\text{C}-\text{C}\equiv\text{C}$ -, $-\text{CO}-\text{S}$ -, $-\text{S}-\text{CO}$ -, $-\text{CO}-\text{O}$ -, $-\text{O}-\text{CO}$ -, a direct single bond when $n\geq 1$,



5

10

15

20

25

30

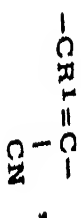
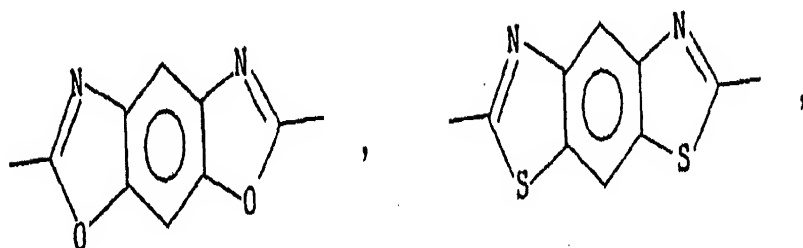
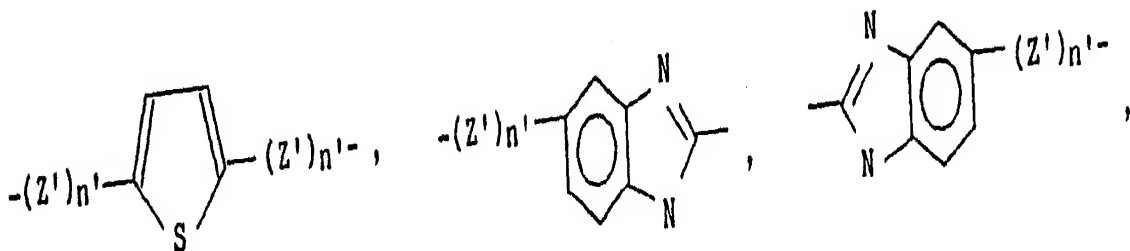
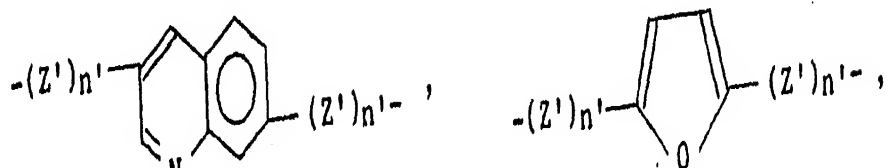
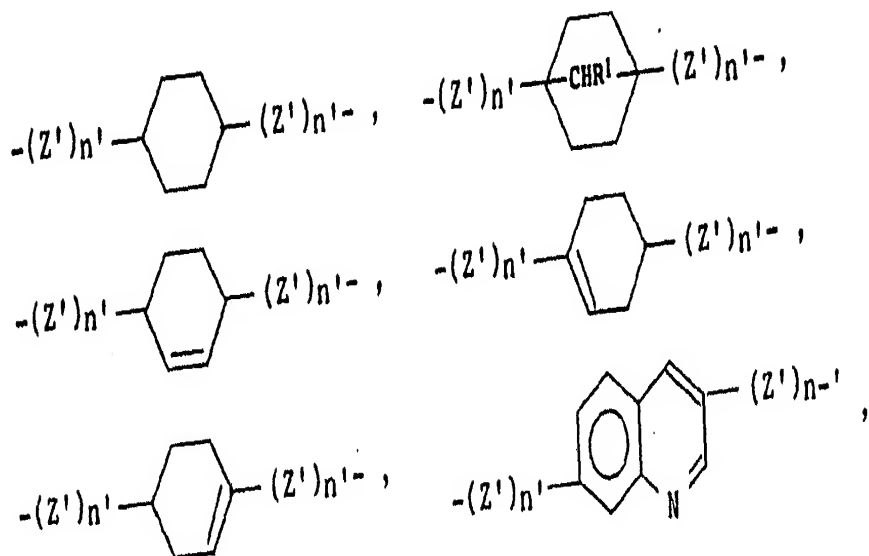
35

40

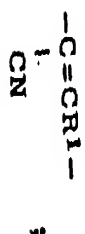
45

50

55

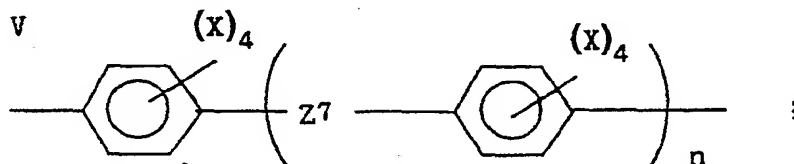


and

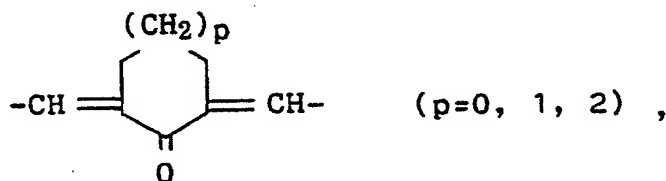
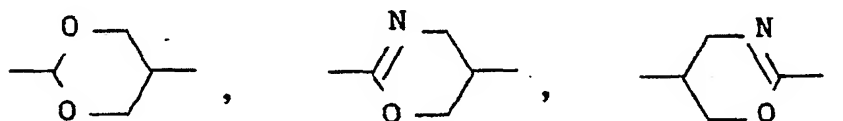
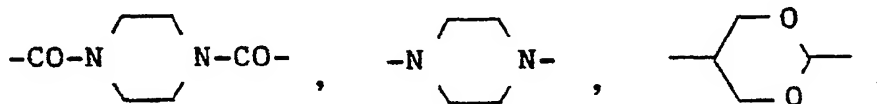
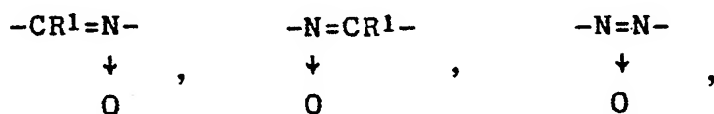


Z' is independently $-O-CO-$, $-CO-O-$, $-NR^1-CO-$, $-CO-NR^1-$; each n' is independently zero or 1; R^1 is independently a hydrogen atom or an alkyl group having from 1 to 4 carbon atoms; Z^6 is a divalent hydrocarbyl group having from 1 to 12, preferably from 1 to 6, more preferably from 1 to 4 carbon atoms; $-SO-$, $-SO_2-$, $-S-$, $-S-S-$, $-O-$, or $-CO-$; each Z^1 is independently $-CO-NH-$, or $-NH-CO-$; Z^2 is a group represented by a cyclic or bicyclic ring system containing from 5 to 12, preferably from 5 to 10, more preferably from 5 to 6 carbon atoms or a group represented by formula V:

Formula V



Z^3 is NH_2- , NH_2-SO_2- , NH_2-CO- , or NH_2-Z^5-O- ; each Z^4 is independently $-CR^1=CR^1-$, $-CR^1=CR^1-$, $-CR^1=N-N=CR^1-$, $-CR^1=CR^1-CO-O-CH_2-$, $-CR^1=CR^1-CO-O-CH_2-CH_2-$, $-CH_2-O-CO-CR^1=CR^1-$, $-CH_2-CH_2-O-CO-CR^1=CR^1-$, $-CR^1=CR^1-CO-O-$, $-O-CO-CR^1=CR^1-$, $-N=N-$, $-CO-NH-$, $-NH-CO-$, $-CO-NH-$, $-NH-CO-$, $-C\equiv C-$, $-C\equiv C-C\equiv C-$, $-CO-S-$, $-S-CO-$, $-CR^1=N-$, $-N=CR^1-$, $-CO-O-$, $-O-CO-$, a direct single bond,



15

20

25

30

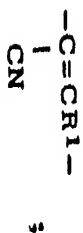
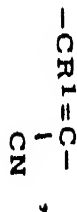
65

40

45

50

55



Z^5 is an alkyl or cycloalkyl group having from 1 to 10, preferably from 1 to 6, more preferably from 1 to 3 carbon atoms; Z^7 is the same as Z^4 with the proviso that it can also independently be a divalent hydrocarbyl group having from 1 to 4 carbon atoms, $-SO-$, $-SO_2-$, $-S-$, $-S-S-$, $-O-$, or $-CO-$; and n has an average value of zero to 6;

5 with the proviso that

(a) at least 80 percent of the Z and X' groups are in the para position with respect to each other in formula I, and

(b) at least 80 percent of the Z^3 and Z^4 groups are in the para position with respect to each other in formula III; and

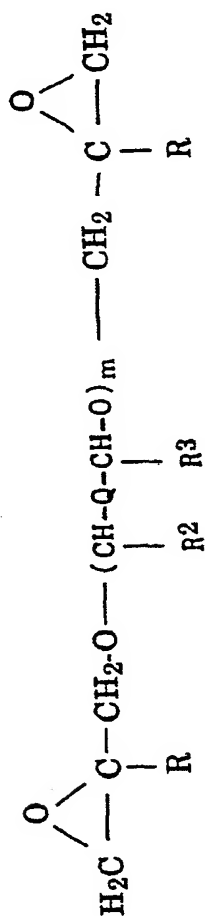
10 wherein components (A) and (B) are employed in amounts which provide a ratio of active hydrogen atoms per vicinal epoxy group of from 0.01:1 to 0.95:1; and

(II) a curing amount of a suitable curing agent or curing catalyst for component (I).

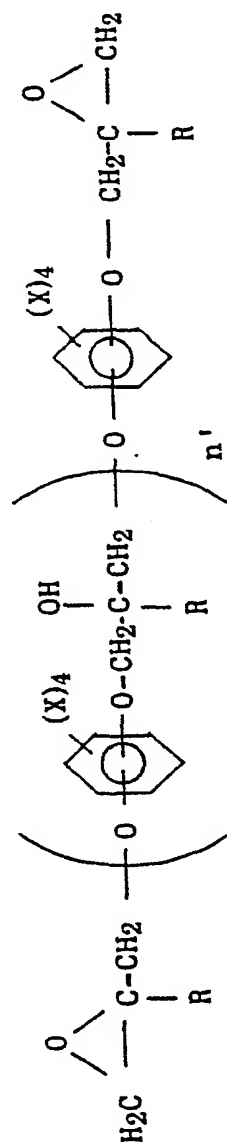
9. A product or article resulting from curing a composition of Claim 8 wherein

(i) component (A) is an epoxy resin represented by the following formulas VI, VII, VIII, IX or X:

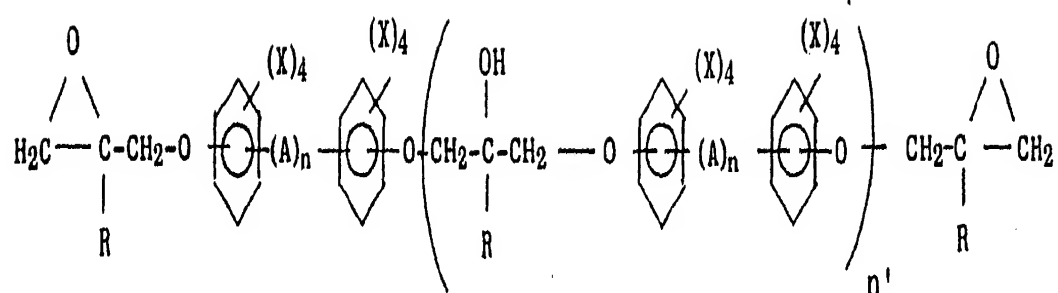
FORMULA VI



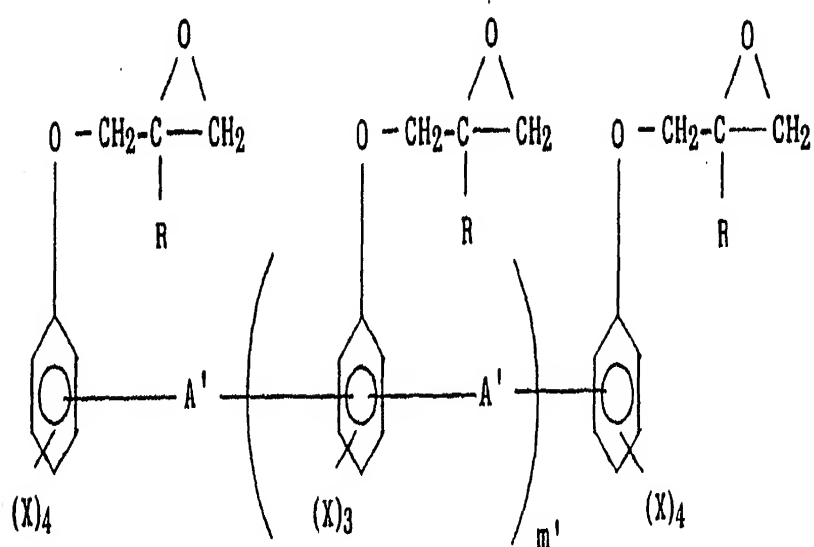
FORMULA VII



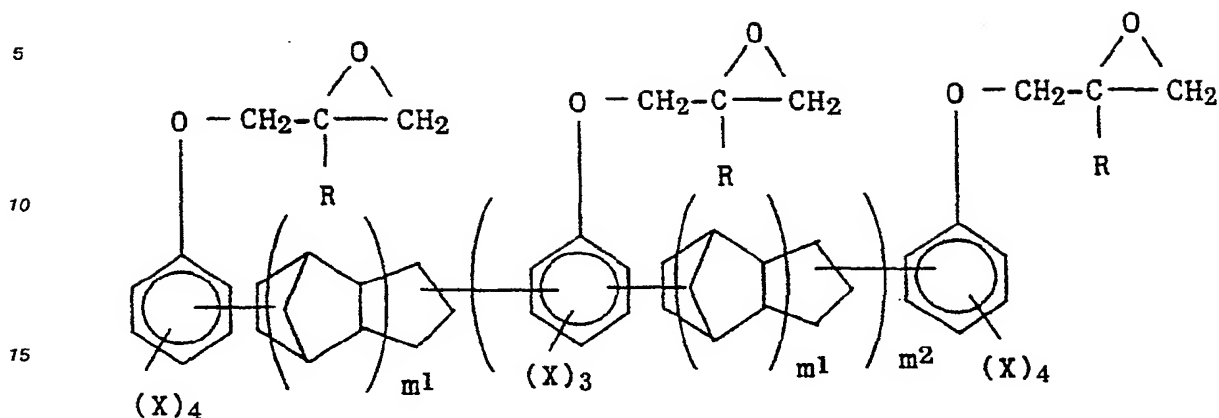
FORMULA VIII



FORMULA IX



FORMULA X

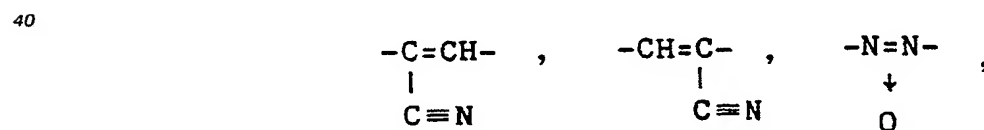


20 wherein each A is independently a divalent hydrocarbonyl group having from 1 to 12 carbon atoms, -O-, -S-, -S-S-, -SO-, -SO₂-, or -CO-; each A' is independently a divalent hydrocarbon group having from 1 to 6, preferably from 1 to 3 carbon atoms; Q is a single bond, -CH₂-S-CH₂-, -(CH₂)_{n1}-, or



each R is independently hydrogen or an alkyl group having from 1 to 4 carbon atoms; each R² and R³ is independently hydrogen, a hydrocarbyl or haloalkyl group having from 1 to 6 carbon atoms; each X is independently hydrogen, a hydrocarbyl or hydrocarbyloxy group having from 1 to 12 carbon atoms, a halogen atom, -NO₂ or -C≡N; m has a value from 1 to 10; m' has a value from 0.01 to 12; m¹ has an average value from 1 to 12; m² has a value from 1 to 12; n has a value of zero or 1; n' has an average value from zero to 3; and n¹ has an average value from 1 to 10;

(ii) component (B) is a compound represented by formula I wherein each X is independently hydrogen, a hydrocarbyl or hydrocarbyloxy group containing from 1 to 4 carbon atoms, a halogen atom or a -C≡N group; each X' is independently a hydroxyl group, a carboxylic acid group or a group represented by formula IV; each Z is independently -N=N-, -CO-NH-, -NH-CO-, -CO-O-, -O-CO-, -CR¹=CR¹,



45 where each R¹ is independently a hydrogen atom or an alkyl group having from 1 to 4 carbon atoms; each Z⁶ is independently a divalent hydrocarbonyl group having from 1 to 4 carbon atoms, -SO-, -SO₂-, -O-, or -CO-; and n has a value from zero to 6;

(iii) components (A) and (B) are employed in amounts which provide a ratio of active hydrogen atoms
50 per vicinal epoxy group of from 0.01:1 to 0.95:1; and

(iv) component (II) is an aliphatic or cycloaliphatic compound containing more than one primary or secondary amine group per molecule, an aromatic compound containing more than one primary or secondary amine group per molecule, dicarboxylic acid anhydride, a phenol-aldehyde novolac resin, a substituted phenol-aldehyde novolac resin, an alkylolated urea-aldehyde resin, an alkylolated melamine-aldehyde resin, a guanadine, or any combination thereof.

10. A product or article resulting from curing a composition of Claim 8 wherein component (B) is bis(4-hydroxyphenyl)terephthalate.

11. A product or article resulting from curing a composition of Claim 9 wherein component (B) is N,N'-

diphenylterephthalamide or a mixture of N,N'-diphenylterephthalamide and N-phenylbenzamide.

12. A product or article resulting from curing a composition of claim 9 wherein component (B) is 4'-sulfonamido-N-phenylbenzamide.

13. The product or article of Claim 8, 9, 10, 11 or 12 wherein said epoxy resin is oriented prior to or during curing.

14. The product or article of Claim 13 wherein said orientation is achieved by subjecting the composition to an electric field, a magnetic field or a shear flow.

10

15

20

25

30

35

40

45

50

55

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) Publication number:

0 379 055 A3

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: **90100485.3**

(51) Int. Cl.⁵: **C08G 59/02, C08G 59/20**

(22) Date of filing: **11.01.90**

(30) Priority: **17.01.89 US 297896**

(43) Date of publication of application:
25.07.90 Bulletin 90/30

(84) Designated Contracting States:
AT BE CH DE ES FR GB GR IT LI NL SE

(68) Date of deferred publication of the search report:
27.05.92 Bulletin 92/22

(71) Applicant: **THE DOW CHEMICAL COMPANY**
2030 Dow Center Abbott Road P.O. Box 1967
Midland Michigan 48640-1967(US)

(72) Inventor: **Earls, Jimmy D.**
209 Banyan
Lake Jackson Texas 77566(US)
Inventor: **Puckett, Paul M.**
126 Daisy
Lake Jackson Texas 77566(US)

(74) Representative: **Sternagel, Hans-Günther, Dr.**
et al
Patentanwälte Dr. Michael Hann Dr. H.-G.
Sternagel Sander Aue 30
W-5060 Bergisch Gladbach 2(DE)

(54) **Mesogenic advanced epoxy compounds.**

(57) Advanced epoxy resin compounds are prepared by reacting weight epoxy resins with active hydrogen-containing compounds which contain rigid, mesogenic type structures.

EP 0 379 055 A3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 90 10 0485

Page 1

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
X	WORLD PATENTS INDEX Derwent Publications Ltd., London, GB; AN 78-59720A & JP-A-53 079 998 (NIPPON TELEG & TELEPH) 14 July 1978	1-14	C08G59/02 C08G59/20
Y	* abstract *	1-14	
Y	EP-A-0 252 359 (BAYER AG) * the whole document *	1-14	
A	US-A-3 182 074 (D. F. LOUCRINI) * column 1, line 38 - column 2, line 5 * * column 6, line 38 - line 41 *	1	
A	MACROMOLECULES vol. 20, no. 8, August 1987, WASHINGTON DC pages 2010 - 2017; SHAUL M. AHORANI: '*Monodisperse Rodlike Oligomers and Their Mesomorphic Higher Molecular Weight Homologues*'		TECHNICAL FIELDS SEARCHED (Int. Cl.5)
A	THE JOURNAL OF ORGANIC CHEMISTRY vol. 41, no. 15, 23 July 1976, EASTON pages 2566 - 2571; D.C. SCHROEDER, J.P. SCHROEDER: 'Liquid Crystals. Mesomorphic Phenols and Primary Amines. p-Phenylene Dibenzoates with Terminal Hydroxy and Amino Groups.'	1	C08G C07D G02F
A	CHEMICAL ABSTRACTS, vol. 103, no. 16, 21 October 1985, Columbus, Ohio, US; abstract no. 124509N, '*Curing agents for epoxy resins*' * abstract *	1	
A	US-A-3 725 341 (M.G. ROGERS, E.A.) * column 3, line 1 - column 4, line 66; claims *	1	
-/-			
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 30 MARCH 1992	Examiner DERAEDT G.
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons * : member of the same patent family, corresponding document	

EPO FORM 1500 01.82 (P.0001)



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 90 10 0485

Page 2

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
A	EP-A-0 287 233 (NORTH DAKOTA STATE UNIVERSITY) * claims * -----	1	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 30 MARCH 1992	Examiner DERAEDT G.
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons ----- & : member of the same patent family, corresponding document			

EPO FORM 1503 03.92 (P0601)